

STIC Search Report Biotech-Chem Library

STIC Database Tracking Number: 139521

TO: Dave Nguyen

Location: REM/2D31/2C18

Art Unit: 1632

Monday, December 13, 2004

Case Serial Number: 10/068160

From: Noble Jarrell

Location: Biotech-Chem Library

Rem 1B71

Phone: 272-2556

Noble.jarrell@uspto.gov

| Search Notes | |
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STIC-Biotech/ChemLib

139501

CRFE

From:

Nguyen, Dave

Sent:

Friday, December 03, 2004 3:50 PM

To: Subject:

STIC-Biotech/ChemLib formula search 10/068160

Please serach the following claim:

A substantially pure or isolated oligodeoxynucleotide of between 18 and 30 nucleotides in length comprising a sequence represented by the following formula:

5' $X_1X_2X_3Pu_1Py_2CpG Pu_3 Py_4X_4X_5X_6(W)_M(G)_N-3'$

wherein the central CpG motif is unmethylated, Pu is a purine nucleotide, Py is a pyrimidine nucleotide, X and W are any nucleotide, M is any integer from 0 to 10, and N is any integer from 4 to 10, and

wherein $X_1X_2X_3Pu_1Py_2$ and Pu_3 $Py_4X_4X_5X_6$ are complementary

MailBox:2c18
Dave Trong Nguyen
Primary Examiner
Remsen 2D31
571-272-0731

DEC -3 2005

STAFF USE ONLY

Searcher: Noble

Searcher Phone: 2Date Searcher Picked up: 17 13 64

Date Completed: 17 13 64

Searcher Prep/Rev. Time: 10

Online Time: 20

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(FILE 'HOME' ENTERED AT 08:46:58 ON 13 DEC 2004)

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FILE 'REGISTRY' ENTERED AT 08:49:20 ON 13 DEC 2004

FILE 'HCAPLUS' ENTERED AT 08:49:22 ON 13 DEC 2004
L5 TRA L4 1- RN : 251 TERMS

FIGHS 'REGISHRY' ENTERED AT 08:49:23 ON 13 DEC 2004

FILE 'HCAPLUS' ENTERED AT 08:50:45 ON 13 DEC 2004

EDGE UHCAPLUS' ENTERED AT 08:51:05 ON 13 DEC 2004
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FILE COVERS 1907 - 13 Dec 2004 VOL 141 ISS 25 FILE LAST UPDATED: 12 Dec 2004 (20041212/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

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ANSWER 1 OF 2 HCAPLUS COPYRIGHT 2004 ACS on STN L4 AN 2003:241991 HCAPLUS DN 138:270283 Entered STN: 28 Mar 2003 Oligodeoxynucleotide and its use to induce an immune response TI IN Klinman, Dennis; Verthelyi, Daniela; Ishii, Ken; Mond, James J.; Gursel, USA U.S. Pat. Appl. Publ., 52 pp., Cont.-in-part of U.S. Ser. No. 958,713. so CODEN: USXXCO DT Patent LΑ English A61K048-00; C07H021-04 IC 514044000; 536023100 NCL CC 15-2 (Immunochemistry) Section cross-reference(s): 1 FAN.CNT 2 PATENT NO. KIND DATE APPLICATION NO. DATE 2003060440 DIS 2002-68166 ΡI A1 20030327 20020206 <--

Search done by Noble Jarrell

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PRAI TS 1999-128898B
                                 20011011
    US 2001-958713
CLASS
PATENT NO.
                 CLASS PATENT FAMILY CLASSIFICATION CODES
                 IC
                        A61K048-00IC
                                          C07H021-04
US 2003060440
                        514044000; 536023100
                 NCL
                 ECLA
                        A61K031/7088; A61K031/7088+M; A61K039/39+M
US 2003060440
    D type CpG oligodeoxynucleotides are provided herein that include a
     sequence represented by the following formula: 5'-X1X2X3Pu1Py2CpGPu3Py4X4X5X6(W)M(G)N-3' wherein the central CpG motif is
     unmethylated, Pu is a purine nucleotide, Py is a pyrimidine nucleotide, X
     and W are any nucleotide, M is any integer from 0 to 10, and N is any
     integer from 4 to 10. The oligodeoxynucleotides can activate immune
     cells, such as antigen-presenting cells or natural killer cell, and/or can
     stimulate production of cytokines. Methods of using these
     oligodeoxynucleotides to induce an immune response are provided. The
     oligodeoxynucleotides can be used in treatment or amelioration of cancer,
     allergy, autoimmune disease, immunodeficiency, or infection. They can
     also be used to enhance the efficacy of vaccines.
     CpG oligodeoxynucleotide immunotherapy immunostimulation
    Allergy
     Allergy inhibitors
     Anti-infective agents
     Antibiotics
     Antitumor agents
     Antiviral agents
     Asthma
     Autoimmune disease
     Cell activation
     Chemotherapy
     Dendritic cell
     Fungicides
     Human
     Immunodeficiency
     Immunostimulation
     Immunotherapy
     Leishmania major
     Monocyte
     Mycosis
     Neoplasm
     Radiotherapy
     Vaccines
        (CpG oligodeoxynucleotides for immunostimulation and immunotherapy of
        various diseases)
     RL: ADV (Adverse effect, including toxicity); BIOL (Biological study)
        (CpG oligodeoxynucleotides for immunostimulation and immunotherapy of
        various diseases)
IT
     Cytokines
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (CpG oligodeoxynucleotides for immunostimulation and immunotherapy of
        various diseases)
IT
     Interleukin 10
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (CpG oligodeoxynucleotides for immunostimulation and immunotherapy of
        various diseases)
IT
     Oligodeoxyribonucleotides
     RL: BSU (Biological study, unclassified); PAC (Pharmacological activity);
     PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES
     (Uses)
        (CpG-containing; CpG oligodeoxynucleotides for immunostimulation and
        immunotherapy of various diseases)
IT
     Immunostimulants
        (adjuvants; CpG oligodeoxynucleotides for immunostimulation and
        immunotherapy of various diseases)
IT
     Infection
        (bacterial; CpG oligodeoxynucleotides for immunostimulation and
        immunotherapy of various diseases)
IT
     Immunity
        (cell-mediated; CpG oligodeoxynucleotides for immunostimulation and
        immunotherapy of various diseases)
     Lipids, biological studies
     RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (delivery complex; CpG oligodeoxynucleotides for immunostimulation and
```

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immunotherapy of various diseases)
IT
     Chemokines
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (interferon .gamma.-inducible protein-10; CpG oligodeoxynucleotides for
        immunostimulation and immunotherapy of various diseases)
     Drug delivery systems
IT
        (liposomes; CpG oligodeoxynucleotides for immunostimulation and
        immunotherapy of various diseases)
IT
    Lymphocyte
        (natural killer cell; CpG oligodeoxynucleotides for immunostimulation
        and immunotherapy of various diseases)
IT
        (solid; CpG oligodeoxynucleotides for immunostimulation and
        immunotherapy of various diseases)
IT
     Infection
        (viral; CpG oligodeoxynucleotides for immunostimulation and
        immunotherapy of various diseases)
    Drug delivery systems
IT
        (virosomes; CpG oligodeoxynucleotides for immunostimulation and
        immunotherapy of various diseases)
IT
     Interferons
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (.alpha.; CpG oligodeoxynucleotides for immunostimulation and
        immunotherapy of various diseases)
IT
     Interferons
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (.gamma.; CpG oligodeoxynucleotides for immunostimulation and
        immunotherapy of various diseases)
TT
     503572-62-1
                   503572-63-2
                                 503572-64-3
                                                503572-65-4
                                                              503572-66-5
                   503572-68-7
                                                503572-70-1
                                                              503572-71-2
     503572-67-6
                                 503572-69-8
                   503572-73-4
     503572-72-3
                                 503572-74-5
                                                503572-75-6
                                                              503572-76-7
     503572-77-8
                   503572-78-9
     RL: BSU (Biological study, unclassified); PAC (Pharmacological activity);
     PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES
     (Uses)
        (CpG oligodeoxynucleotides for immunostimulation and immunotherapy of
        various diseases)
IT
     503540-93-0
                  503540-94-1
     RL: PRP (Properties)
        (Unclaimed; oligodeoxynucleotide and its use to induce an immune
        response)
IT
     57-88-5, Cholesterol, biological studies
     RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (delivery complex; CpG oligodeoxynucleotides for immunostimulation and
        immunotherapy of various diseases)
     503575-68-6
                   503575-69-7
IT
                                 503575-70-0
                                                503575-71-1
                                                              503575-72-2
     503575-73-3
                                 503575-75-5
                   503575-74-4
                                                503575-76-6
                                                              503575-77-7
     503575-78-8
                   503575-79-9
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                   503575-84-6
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                                                503575-86-8
                                                              503575-87-9
     503575-88-0
                   503575-89-1
                                 503575-90-4
                                                503575-91-5
                                                              503575-92-6
     503575-93-7
                   503575-94-8
                                 503575-95-9
                                                503575-96-0
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                  503576-04-3
                                                              503576-07-6
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                   503576-14-5
                                 503576-15-6
                                                503576-16-7
                                                              503576-17-8
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                                                503576-51-0
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                   503576-54-3
                                                503576-56-5
                                                              503576-57-6
                                 503576-55-4
     503576-58-7
                   503576-59-8
     RL: PRP (Properties)
        (unclaimed nucleotide sequence; oligodeoxynucleotide and its use to
        induce an immune response)
IT
     503576-60-1
                   503576-61-2
                                 503576-62-3
                                                503576-63-4
                                                              503576-64-5
     503576-65-6
                   503576-66-7
     RL: PRP (Properties)
        (unclaimed sequence; oligodeoxynucleotide and its use to induce an
        immune response)
     ANSWER 2 OF 2 HCAPLUS COPYRIGHT 2004 ACS on STN
L4
     2000:741930 HCAPLUS
AN
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133:320986
ED
     Entered STN: 20 Oct 2000
     Oligodeoxynucleotide and its use to induce an immune response
TΙ
     Klinman, Dennis; Ishii, Ken; Verthelyi, Daniela
PA
     United States Dept. of Health and Human Services, USA
     PCT Int. Appl., 46 pp.
SO
     CODEN: PIXXD2
DT
     Patent
LA
     English
     ICM A61K031-70
IC
     ICS A61K039-39; C07H021-00
CC
     15-2 (Immunochemistry)
     Section cross-reference(s): 3, 63
FAN.CNT 2
     PATENT NO.
                          KIND
                                  DATE
                                               APPLICATION NO.
                                                                        DATE
                           ----
     A2
                                  20001019
                                                                        20000412 <--
PΙ
     WO 2000061151
                           A3
                                  20010426
         W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR,
             CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU,
             ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU,
             LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE,
             SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
         RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE,
             DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF,
                                 , GW, ML, MR, NE, SN, TD, TG
20020206 EP 2000-923283
             CG, CI, CM, GA, GN, GW, ML, MR, NE,
                                                             TG
                           200000412 <=-
     EP 11.76966
                  BE,
                      CĤ,
                          DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, SI, LT, LV, FI, RO
PRAI US 1999-128898P
                                  19990412 <--
                           P
     WO 2000-US9839
                           W
                                  20000412 <--
CLASS
 PATENT NO.
                  CLASS PATENT FAMILY CLASSIFICATION CODES
 WO 2000061151
                  ICM
                         A61K031-70
                         A61K039-39; C07H021-00
                  ICS
     The present invention provides a substantially pure or isolated
     oligodeoxynucleotide of at least about 10 nucleotides comprising a
     sequence represented by either the formula: 5' N1N2N3T-CpG-WN4N5N6 3'
     wherein the central CpG motif is unmethylated, W is A or T, and N1, N2,
     N3, N4, N5, and N6 are any nucleotides, or the formula: 5' RY-CpG-RY 3'
     wherein the central CpG motif is unmethylated, R is A or G, and Y is C or
     T, as well as an oligodeoxynucleotide delivery complex and a pharmacol. composition comprising the present inventive oligodeoxynucleotide, and a method
     of inducing an immune response by administering the present inventive oligodeoxynucleotide to a host. The oligodeoxynucleotides with phosphate
     or phosphorothicate backbone modification are useful for inducing
     cell-mediated and humoral immune response and are therefore useful for
     treatment of allergy, asthma, cancer, autoimmune disease, immunol.
     disease, infection, and immune deficiency.
     oligodeoxynucleotide adjuvant vaccine allergy cancer infection; autoimmune
     disease asthma oligodeoxynucleotide adjuvant vaccine
IT
     Cell activation
         (B cell; oligodeoxynucleotide for use to induce immune response)
     B cell (lymphocyte)
IT
     Lymphocyte
         (activation; oligodeoxynucleotide for use to induce immune response)
TT
     Infection
        (anti-; oligodeoxynucleotide for use to induce immune response)
IT
     Chemical warfare agents
        (bio-; oligodeoxynucleotide for use to induce immune response)
IT
     Diagnosis
        (cancer; oligodeoxynucleotide for use to induce immune response)
IT
     Drug delivery systems
        (carriers; oligodeoxynucleotide for use to induce immune response)
     Immunity
IT
         (cell-mediated; oligodeoxynucleotide for use to induce immune response)
IT
        (diagnosis; oligodeoxynucleotide for use to induce immune response)
IT
     Immunity
         (disorder; oligodeoxynucleotide for use to induce immune response)
IT
     Immunity
        (humoral; oligodeoxynucleotide for use to induce immune response)
TT
     Drug delivery systems
        (liposomes; oligodeoxynucleotide for use to induce immune response)
```

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ΙT
    Cell activation
        (lymphocyte; oligodeoxynucleotide for use to induce immune response)
TΤ
        mal cell
        (non-B; oligodeoxynucleotide for use to induce immune response)
IT
    Allergy
    Allergy inhibitors
    Antitumor agents
    Asthma
    Autoimmune disease
    B cell (lymphocyte)
    Drug targeting
     Immunodeficiency
     Immunostimulants
    Neoplasm
    Vaccines
        (oligodeoxynucleotide for use to induce immune response)
    Antibodies
     Cytokines
    Interleukin 6
    RL: BOC (Biological occurrence); BSU (Biological study, unclassified); MFM
     (Metabolic formation); THU (Therapeutic use); BIOL (Biological study);
    FORM (Formation, nonpreparative); OCCU (Occurrence); USES (Uses)
        (oligodeoxynucleotide for use to induce immune response)
    Lipids, biological studies
    Oligodeoxyribonucleotides
    RL: BSU (Biological study, unclassified); PRP (Properties); THU
     (Therapeutic use); BIOL (Biological study); USES (Uses)
        (oligodeoxynucleotide for use to induce immune response)
IT
    Neoplasm
        (solid; oligodeoxynucleotide for use to induce immune response)
IT
    Antisense oligonucleotides
    RL: BOC (Biological occurrence); BSU (Biological study, unclassified); MFM
     (Metabolic formation); THU (Therapeutic use); BIOL (Biological study);
    FORM (Formation, nonpreparative); OCCU (Occurrence); USES (Uses)
        (therapy; oligodeoxynucleotide for use to induce immune response)
    Liposomes
IT
        (virosomes; oligodeoxynucleotide for use to induce immune response)
IT
    RL: BOC (Biological occurrence); BSU (Biological study, unclassified); MFM
     (Metabolic formation); THU (Therapeutic use); BIOL (Biological study);
    FORM (Formation, nonpreparative); OCCU (Occurrence); USES (Uses)
        (.gamma.; oligodeoxynucleotide for use to induce immune response)
    199810-73-6D, phosphate or phosphorothicate derivs.
                                                           250141-69-6D,
                               oate derivs. 301939-29-7D, phosphate or 301939-30-0D, phosphate or phosphorothioate
    phosphate or phosphorothicate derivs.
    phosphorothioate derivs.
              301939-31-1D, phosphate or phosphorothioate derivs.
    301939-32-2D, phosphate or phosphorothicate derivs.
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                                                           301939-38-8D,
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    phosphorothicate derivs.
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              301939-46-8D, phosphate or phosphorothicate derivs.
    301939-47-9D, phosphate or phosphorothicate derivs.
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              301939-51-5D, phosphate or phosphorothicate derivs.
    derivs.
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                               301939-60-6D, phosphate or phosphorothioate
    phosphorothicate derivs.
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    301939-62-8D, phosphate or phosphorothicate derivs.
                                                           301939-63-9D
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                               301939-65-1D, phosphate or phosphorothioate
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              301939-66-2D, phosphate or phosphorothioate derivs.
    derivs.
    301939-67-3D, phosphate or phosphorothicate derivs.
                                                            301939-68-4D,
    phosphate or phosphorothicate derivs.
                                             301939-69-5D, phosphate or
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<<<

RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (oligodeoxynucleotide for use to induce immune response)

=> b wpix

FIGE ** WPIX** ENTERED AT 08:51:14 ON 13 DEC 2004

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L10 ANSWER 1 OF 1 WPIX COPYRIGHT 2004 THE THOMSON CORP on STN

AN 2001-006880 [01] WPIX

DNC C2001-001559

TI Novel oligonucleotides useful for the prevention and treatment of allergies, cancer, and autoimmune disorders and for ameliorating symptoms resulting from exposure to a bio-warfare agent.

DC 804 D16

IN ISHII, K; KLINMAN, D; VERTHELYI, D; GURSEL, M; MOND, J J

PA (USSH) US DEPT HEALTH & HUMAN SERVICES; (GURS-I) GURSEL M; (ISHI-I) ISHII K; (KLIN-I) KLINMAN D; (MOND-I) MOND J J; (VERT-I) VERTHELYI D

CYC 93

PI WO 2000061151 A2 20001019 (200101)* EN 46 A61K031-70 RW: AT BE CH CY DE DK EA ES FI FR GB GH GM GR IE IT KE LS LU MC MW NL OA PT SD SE SL SZ TZ UG ZW

W: AE AG AL AM AT AU AZ BA BB BG BR BY CA CH CN CR CU CZ DE DK DM DZ EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MA MD MG MK MN MW MX NO NZ PL PT RO RU SD SE SG SI

SK SL TJ TM TR TT TZ UA UG US UZ VN YU ZA ZW

AU 2000043437 A 20001114 (200108) A61K031-70 EP 1176966 A2 20020206 (200218) EN A61K031-70

R: AL AT BE CH CY DE DK ES FI FR GB GR IE IT LI LT LU LV MC MK NL PT RO SE SI

US 2003060440 A1 20030327 (200325) A61K048-00 <
ADT WO 2000061151 A2 WO 20000-US9839 20000412; AU 2000043437 A AU 2000-43437 20000412; EP 1176966 A2 EP 2000-923283 20000412, WG 20000-US9839 20000412; US 2003060440 A1 Provisional US

2000-059889 2000-05927 US 2003060440 A1 (<u>Provisional</u>) US <u>ମ୍ୟୁତ୍ୟ ମଧ୍ୟ ପ୍ରଥମ ଅନ୍ତର୍ଗ ମଧ୍ୟ ପ୍ରଥମ</u> CIP of US 2001-958713 20011010, US 2002-68160 20020206

FDT AU 2000043437 A Based on WO 2000061151; EP 1176966 A2 Based on WO 2000061151

PRAI 0S 1959-128898P 19990412; US 2001-958713 20011010; US 2002-68160 20020206

IC ICM A61K031-70; A61K048-00

ICS A61K039-39; C07H021-00; C07H021-04

AB WO 200061151 A UPAB: 20001230

NOVELTY - A pure or isolated oligonucleotide (I) of at least 10 nucleotides, comprising a specific sequence, is new.

DETAILED DESCRIPTION - A pure or isolated oligonucleotide (I) of at least 10 nucleotides, comprising a specific sequence, is new. (I)

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comprises the sequence: 5'-N1N2N3T-CpG-WN4N5N6-3' or 5'-RY-CpG-RY-3'.
     W = A \text{ or } T;
          N1-6 = any nucleotides:
     R = A \text{ or } G; \text{ and }
     Y = C \text{ or } T.
           The central CpG motif is unmethylated.
           INDEPENDENT CLAIMS are also included for the following:
           (1) an oligonucleotide delivery complex (II), comprising (I) and a
     targeting agent; and
           (2) a pharmacological composition comprising (I).
           ACTIVITY - Cytostatic; Immunosuppressive; Antiarthritic;
     Antimicrobial; Antiallergic; Anti-HIV (human immunodeficiency virus);
     Protozoacide; Tuberculostatic; Antiasthmatic; Dermatological;
     Neuroprotective. No biological data is given.
           MECHANISM OF ACTION - Inducer of cell or humoral mediated immune
     response. Vaccine. Oligodeoxynucleotides (ODNs) ATCGACTCTCGAGCGTTCT and
     ATCGACTAGCGTTCGTTCTC were synthesized on a DNA synthesizer and the
     induction of immune response by the ODNs was tested. The normal DNA
     backbone phosphodiesterase linkages were replaced with phosphorothioate
     linkages. Human peripheral blood mononuclear cells were incubated for 72
     hours with the ODNs. Interleukin (IL-6) and tumor necrosis factor gamma
     (TNF- gamma ) levels were determined by enzyme linked immunosorbent assay
     (ELISA) using anti-IL-6 and anti-TNF- gamma antibodies. Cell proliferation
     was determined by (3H) thymidine incorporation. The results showed that
     the sequence containing 5'-N1N2N3T-CpG-WN4N5N6 3' was desirable to induce
     a humoral immune response. In addition, maximum induction was observed for
     ODNs that contained a phosphorothioate backbone. The sequence containing
     5'RY-CpG-RY 3', was desirable to induce a cell-mediated immune response.
     The IL-6 levels and cell proliferation obtained with ATCGACTCTCGAGCGTTCT
     was 85 and 44 and the IL-6 level obtained with ATCGACTAGCGTTCGTTCTC was
          USE - (I) is useful for inducing cell-mediated immune response in
     humans by activating non-B cells or inducing humoral-mediated immune
     response by activating B cells in the human. (I) induces cytokine e.g.
     interleukin (IL)-6, or antibody production in the host. The induction of
     immune response is useful for treating, preventing or ameliorating an allergic reaction, preferably asthma or an infection, where (I) is
     administered either alone or in combination with an anti-allergenic agent
     or anti-infectious agent, autoimmune disorder, a disease associated with
     immune system deficiency and symptoms resulting from exposure to a
     bio-warfare agent. (I) alone or in combination with an anti-cancer agent is useful for treating solid tumor cancer. The induction of immune
     response is used in antisense therapy and to improve the efficacy of a
     vaccine. (I) is preferably administered to lymphocytes ex vivo, producing
     activated lymphocytes which are then administered to the host (claimed).
     Allergic conditions include eczema, allergic rhinitis, food allergies and
     other atopic conditions, immune system deficiencies e.g. lupus
     erythematosus and autoimmune diseases include rheumatoid arthritis and
     multiple sclerosis and infections include viral, bacterial and fungi infections such as tuberculosis, acquired immunodeficiency syndrome
     (AIDS), leishmania and schistosomiasis.
          ADVANTAGE - The oligodeoxynucleotides are more resistant to
     degradation.
     Dwg.0/0
     CPT
     AB; DCN
     CPI: B01-D02; B04-E02F; B04-E06; B14-A01; B14-A02; B14-A02B1; B14-A04;
          B14-C06; B14-C09B; B14-G01B; B14-G02A; B14-G02D; B14-H01; B14-H01B;
          B14-K01A; B14-N17C; B14-S01; B14-S03; B14-S11; D05-H07; D05-H12B2
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FA MC

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L4
               2 L1-3
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                TRA L4 1- RN :
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L8
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L10
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              61 L24 AND P/DT
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             25 1/25 AND DE/DC B
127
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substance identification.

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ANSWER 1 OF 21 HCAPLUS COPYRIGHT 2004 ACS on STN
L21
     2004:995900 HCAPLUS
AN
     Entered STN: 19 Nov 2004
ED
     Method of preventing infections from bioterrorism agents with
     immunost imedatory CPG oligonucleotides
[Minman, Dennis W., Ivins, Bruce; Westhelmin Denniela
TN
     The Government of the United States of America as Represented by the
     Secretary of the Department of Health and Human
     Services, USA
     PCT Int. Appl., 142 pp.
SO
     CODEN: PIXXD2
DT
     Patent
     English
LA
IC
     ICM A61K
CC
     1-5 (Pharmacology)
     Section cross-reference(s): 3
FAN.CNT 1
     PATENT NO.
                          KIND
                                 DATE
                                              APPLICATION NO.
                                                                       DATE
     WO 2004098491
                           A2
                                 20041118
                                             WO 2003-US34523
                                                                       20031031
         W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH,
             CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD,
             GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,
             LK, LR, LS, LT, LU, LV, MA, MD, MG, MK; MN, MW, MX, MZ, NI, NO,
             NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ,
         TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ,
             BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE,
             ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK,
             TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
PRAI US 2002-422964P
                           Р
                                 20021101
CLASS
                  CLASS PATENT FAMILY CLASSIFICATION CODES
 PATENT NO.
 WO 2004098491 ICM A61K
     The present disclosure relates to a method of preventing or treating an
     infection caused by a bioterrorism agent, specifically to a method of
     increasing an immune response to a bioterrorism agent using an
     oligodeoxynucleotide including a CpG motif, and a method of enhancing the
     immunogenicity of a vaccine against a bioterrorism agent using an
     oligodeoxynucleotide including a CpG motif.
     infection bioterrorism immunostimulatory CPG oligonucleotide
ST
     G proteins (guanine nucleotide-binding proteins)
IT
     RL: BSU (Biological study, unclassified); BIOL (Biological study) (Rac; method of preventing infections from bioterrorism agents with
        immunostimulatory CPG oligonucleotides)
IT
     Bacillus anthracis
         (anthrax from; method of preventing infections from bioterrorism agents
        with immunostimulatory CPG oligonucleotides)
IT
         (bacterial; method of preventing infections from bioterrorism agents
        with immunostimulatory CPG oligonucleotides)
IT
     Terrorism
         (bioterrorism; method of preventing infections from bioterrorism agents
        with immunostimulatory CPG oligonucleotides)
IT
     Staphylococcus
         (enterotoxin B; method of preventing infections from bioterrorism
        agents with immunostimulatory CPG oligonucleotides)
TT
     Oligodeoxyribonucleotides
     RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
         (immunostimulatory; method of preventing infections from bioterrorism
        agents with immunostimulatory CPG oligonucleotides)
IT
     Drug delivery systems
        (inhalants; method of preventing infections from bioterrorism agents
        with immunostimulatory CPG oligonucleotides)
IT
     Drug delivery systems
         (injections, i.m.; method of preventing infections from bioterrorism
        agents with immunostimulatory CPG oligonucleotides)
TT
     Drug delivery systems
         (injections, i.v.; method of preventing infections from bioterrorism
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agents with immunostimulatory CPG oligonucleotides)
IT
     Drug delivery systems
         (injections, s.c.; method of preventing infections from bioterrorism
        agents with immunostimulatory CPG oligonucleotides)
IT
     Anti-infective agents
     Antibiotics
     Antiviral agents
     Bacillus anthracis
     Combination chemotherapy
     Drug bioavailability
     Ebola virus
     Encephalitis
     Escherichia coli
     Fungicides
     Haemophilus influenzae
     Histoplasma capsulatum
     Immunostimulants
     Salmonella
     Shellfish
     Shigella dysenteriae
     Shigella flexneri
     Tick-borne encephalitis virus
     Vaccines
     Variola major virus
     Venoms
     Yersinia pestis
         (method of preventing infections from bioterrorism agents with
        immunostimulatory CPG oligonucleotides)
IT
     Aflatoxins
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (method of preventing infections from bioterrorism agents with immunostimulatory CPG oligonucleotides)
     Antibodies and Immunoglobulins
IT
     RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
         (method of preventing infections from bioterrorism agents with
        immunostimulatory CPG oligonucleotides)
TT
     Ricins
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (ricin toxin; method of preventing infections from bioterrorism agents
        with immunostimulatory CPG oligonucleotides)
IT
     Encephalitis
         (tick-borne; method of preventing infections from bioterrorism agents
        with immunostimulatory CPG oligonucleotides)
IT
     Mycotoxins
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
         (trichothecene; method of preventing infections from bioterrorism
        agents with immunostimulatory CPG oligonucleotides)
IT
     Infection
         (variola; method of preventing infections from bioterrorism agents with
        immunostimulatory CPG oligonucleotides)
IT
     Infection
         (viral; method of preventing infections from bioterrorism agents with
        immunostimulatory CPG oligonucleotides)
TΫ́
     35523-89-8. Saxitoxin
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     (Biological study); USES (Uses)
        (method of preventing infections from bioterrorism agents with immunostimulatory CPG oligonucleotides)
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     794238-68-9 794238-69-0 794238-70-3
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     794238-97-4 794238-98-5 794238-99-6
                                                 794239-00-2
     RL: PRP (Properties)
        (unclaimed nucleotide sequence; method of preventing infections from
        bioterrorism agents with immunostimulatory CPG oligonucleotides)
     794235-01-1
     RL: BSU (Biological study, unclassified); PAC (Pharmacological activity);
     PKT (Pharmacokinetics); PRP (Properties); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (method of preventing infections from bioterrorism agents with
        immunostimulatory CPG oligonucleotides)
     794235-01-1 HCAPLUS
     DNA, d(N-N-T-G-C-A-T-C-G-A-T-G-C-A-G-G-G-G-G) (9CI) (CA INDEX NAME)
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
L21 ANSWER 2 OF 21 HCAPLUS COPYRIGHT 2004 ACS on STN
     2004:496197 HCAPLUS
     141:275892
     Entered STN: 20 Jun 2004
     CpG oligodeoxynucleotides improve the response to hepatitis B immunization
     in healthy and SIV-infected rhesus macaques
     Verthelyi, Daniela; Wang, Vivian W.; Lifson, Jeffrey D.;
     Klinman, Dennis M.
     Division of Therapeutic Proteins, Center for Biologics Evaluation and
     Research, Food and Drug Administration, Bethesda, USA
     AIDS (London, United Kingdom) (2004), 18(7), 1003-1008
     CODEN: AIDSET; ISSN: 0269-9370
     Lippincott Williams & Wilkins
     Journal
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IT

RN

CN

AN

DN

ΤI

ΑU

CS

SO

PB

DT

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LΑ
     English
CC
     15-2 (Immunochemistry)
     Objective: The development of an immunogenic vaccine against hepatitis B
AR
     virus (HBV) is particularly important for HIV-infected patients since
     shared epidemiol. risks result in HIV-infected subjects having a high
     incidence of HBV, and coinfection with HBV increases the occurrence of
     hepatotoxicity with antiretroviral therapy. Although HBV vaccination is recommended to all HIV-pos. patients, its efficacy in these patients is reduced. Methods: Healthy (n = 15) and SIV-infected (n = 17) rhesus
     macaques were immunized with Engerix B alone or combined with type D or
     type K CpG ODN. SIV plasma RNA levels were determined by a real time reverse
     transcriptase polymerase chain reaction and antibody titers to HBV surface
     antigen (HbsAg) were measured by ELISA every 2 wk. Results: In healthy
     macaques, adding D or K ODN to Engerix B accelerated and boosted the titer
     of the anti-HbsAg response. In SIV-infected macaques, Engerix B alone
     elicited no detectable antibody response but a significant response was
     seen when it was combined with K or D ODN. The antibody titer induced by
     vaccinating HIV-infected macaques was inversely correlated with their
     initial viral load, with animals having >107 copies/mL being unable to
     mount a significant response. No adverse events or changes in SIV viral
     load were evident during the study. Conclusions: These findings support
     the development of clin. studies to assess the use of CpG ODN as an
     adjuvant for HBV vaccination in healthy and immunocompromised HIV-infected
     subjects.
     hepatitis B virus vaccine CpG oligodeoxynucleotide SIV infection macaque
ST
IT
     Vaccines
         (AIDS; CpG oligodeoxynucleotides improve the response to hepatitis B
        immunization in healthy and SIV-infected rhesus macaques)
IT
     AIDS (disease)
     Hepatitis B virus
     Human immunodeficiency virus
     Macaca mulatta
     Simian immunodeficiency virus
     Vaccines
        (CpG oligodeoxynucleotides improve the response to hepatitis B
        immunization in healthy and SIV-infected rhesus macaques)
     Antibodies and Immunoglobulins
TT
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
         (CpG oligodeoxynucleotides improve the response to hepatitis B
        immunization in healthy and SIV-infected rhesus macaques)
     Oligodeoxyribonucleotides
TΤ
     RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (CpG oligodeoxynucleotides improve the response to hepatitis B
        immunization in healthy and SIV-infected rhesus macaques)
TT
     Immunostimulants
         (adjuvants; CpG oligodeoxynucleotides improve the response to hepatitis
        B immunization in healthy and SIV-infected rhesus macaques)
IT
     Antigens
     RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
         (hepatitis B surface; CpG oligodeoxynucleotides improve the response to
        hepatitis B immunization in healthy and SIV-infected rhesus macaques)
IT
     Immunization
        (vaccination; CpG oligodeoxynucleotides improve the response to
        hepatitis B immunization in healthy and SIV-infected rhesus macaques)
IT
     Anti-AIDS agents
         (vaccines; CpG oligodeoxynucleotides improve the response to hepatitis
        B immunization in healthy and SIV-infected rhesus macaques)
IT
     Infection
        (viral; CpG oligodeoxynucleotides improve the response to hepatitis B
        immunization in healthy and SIV-infected rhesus macaques)
     351186-51-1, Engerix B 756902-34-8 756902-35-9
IT
     756902-36-0
                   757252-01-0 757252-02-1
                                                  757252-03-2
     RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (CpG oligodeoxynucleotides improve the response to hepatitis B
        immunization in healthy and SIV-infected rhesus macaques)
              THERE ARE 33 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT 33
RE
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(31) Verthelyi, D; J Immunol 2003, V170, P4717 HCAPLUS
(32) Verthelyi, D; Trends Immunol 2003, V24, P519 HCAPLUS
(33) Wong, E; Int J STD AIDS 1996, V7, P490 MEDLINE
IT
     756902-34-8
     RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
         (CpG oligodeoxynucleotides improve the response to hepatitis B
        immunization in healthy and SIV-infected rhesus macaques)
RN
     756902-34-8 HCAPLUS
CN
     DNA, d(G-sp-G-sp-T-G-C-A-T-C-G-A-T-G-C-A-G-G-sp-G-sp-G-sp-G-sp-G) (9CI)
     (CA INDEX NAME)
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
     ANSWER 3 OF 21 HCAPLUS COPYRIGHT 2004 ACS on STN
L21
     2004:451477 HCAPLUS
AN
DN
     141:33759
ED
     Entered STN: 04 Jun 2004
     Method of treating and preventing infections in immunocompromised subjects
     with immunostimulatory CpG oligonucleotides
     Klinman, Dennis M.; Verthelyi, Daniela
IN
PA
     The Government of the USA as Represented by the Secretary of the Dept. of
     Health & Human Services, USA
     U.S. Pat. Appl. Publ., 64 pp.
     CODEN: USXXCO
DТ
     Patent
LA
     English
IC
     ICM A61K031-70
     ICS A01N043-04; A61K039-395; A61K039-12; C12N005-06; C12N005-16
NCL
     424204100; 424130100; 435345000
CC
     1-5 (Pharmacology)
     Section cross-reference(s): 15
FAN.CNT 1
     PATENT NO.
                                               APPLICATION NO.
                          KIND
                                  DATE
                                                                        DATE
                           ----
     US 2004105872
                           A1
                                  20040603
                                               US 2003-666022
                                                                        20030917
PRAI US 2002-411944P
                                  20020918
CLASS
 PATENT NO.
                  CLASS PATENT FAMILY CLASSIFICATION CODES
 US 2004105872
                  ICM
                         A61K031-70
                  ICS
                         A01N043-04; A61K039-395; A61K039-12; C12N005-06;
                         C12N005-16
                  NCL
                         424204100; 424130100; 435345000
US 2004105872
                 ECLA
                        A61K031/7115; A61K031/7125; A61K039/39
     A method is disclosed herein for increasing an immune response to an
     opportunistic infection in an immunocompromised subject. In one
     embodiment, the subject is infected with a lentivirus. The method
     includes increasing an immune response to a pathogen using D
     oligodeoxynucleotides including a CpG motif. Addition of K or D CpG
     oligodeoxynucleotides boosted the immunogenicity of the Engerix B
     hepatitis B virus vaccine to render refractory SIV-infected macaques
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responsive to vaccination.
ST
     opportunistic infection immunocompromised immunostimulatory CpG
     oligonucleotide; lentivirus infection treatment immunostimulatory CpG
     oligonucleotide; hepatitis B virus vaccine immunostimulatory
     oligodeoxynucleotide SIV
TТ
    Hepatitis
        (B, opportunistic infection; immunostimulatory CpG oligonucleotides for
        treating and preventing infections in immunocompromised subjects)
IT
     Oligodeoxyribonucleotides
     RL: BSU (Biological study, unclassified); PAC (Pharmacological activity);
     PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES
     (Uses)
        (D or K, having CpG motif; immunostimulatory CpG oligonucleotides for
        treating and preventing infections in immunocompromised subjects)
TT
        (Kaposi's, opportunistic; immunostimulatory CpG oligonucleotides for
        treating and preventing infections in immunocompromised subjects)
IT
     Vaccines
        (adjuvants for; immunostimulatory CpG oligonucleotides for treating and preventing infections in immunocompromised subjects)
     Immunostimulants
IT
        (adjuvants; immunostimulatory CpG oligonucleotides for treating and
        preventing infections in immunocompromised subjects)
IT
     Mycosis
        (as opportunistic infection; immunostimulatory CpG oligonucleotides for
        treating and preventing infections in immunocompromised subjects)
IT
     Aspergillus
        (aspergillosis from; immunostimulatory CpG oligonucleotides for
        treating and preventing infections in immunocompromised subjects)
IT
        (bacillary angiomatosis, opportunistic; immunostimulatory CpG
        oligonucleotides for treating and preventing infections in
        immunocompromised subjects)
IT
     Infection
        (bacterial, as opportunistic infection; immunostimulatory CpG
        oligonucleotides for treating and preventing infections in
        immunocompromised subjects)
IT
     Candida
        (candidiasis from, opportunistic; immunostimulatory CpG
        oligonucleotides for treating and preventing infections in
        immunocompromised subjects)
IT
     Nervous system, neoplasm
        (central, primary lymphoma, opportunistic; immunostimulatory CpG
        oligonucleotides for treating and preventing infections in
        immunocompromised subjects)
     Granulomatous disease
TΤ
        (chronic, PBMC from humans with, response to CpG oligodeoxynucleotides;
        immunostimulatory CpG oligonucleotides for treating and preventing
        infections in immunocompromised subjects)
IT
     Mycosis
        (coccidioidomycosis, opportunistic; immunostimulatory CpG
        oligonucleotides for treating and preventing infections in
        immunocompromised subjects)
IT
     Immunity
        (disorder, immunocompromised; immunostimulatory CpG oligonucleotides
        for treating and preventing infections in immunocompromised subjects)
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (enhancement of production of; immunostimulatory CpG oligonucleotides for
        treating and preventing infections in immunocompromised subjects)
IT
     Cell proliferation
        (enhancement of; immunostimulatory CpG oligonucleotides for treating
        and preventing infections in immunocompromised subjects)
IT
    Mouth, disease
        (hairy leukoplakia, opportunistic; immunostimulatory CpG
        oligonucleotides for treating and preventing infections in
        immunocompromised subjects)
IT
        (hepatitis B, adjuvants for; immunostimulatory CpG oligonucleotides for
        treating and preventing infections in immunocompromised subjects)
    Antiviral agents
        (highly active antiretroviral therapy (HAART) combination;
        immunostimulatory CpG oligonucleotides for treating and preventing
        infections in immunocompromised subjects)
     Human immunodeficiency virus
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Human immunodeficiency virus 1

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Human immunodeficiency virus 2
    Simian immunodeficiency virus
        (immunocompromisation from infection with; immunostimulatory CpG
        oligonucleotides for treating and preventing infections in
        immunocompromised subjects)
TΤ
        (immunoprotective CpG oligodeoxynucleotides; immunostimulatory CpG
        oligonucleotides for treating and preventing infections in
        immunocompromised subjects)
    AIDS (disease)
    Human
    Immunity
    Immunostimulants
    Infection
    Leishmania amazonensis
    Leishmania major
    Tuberculosis
        (immunostimulatory CpG oligonucleotides for treating and preventing
        infections in immunocompromised subjects)
IT
    Phosphorothioate oligodeoxyribonucleotides
    RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
     (Biological study)
        (immunostimulatory oligodeoxynucleotides containing; immunostimulatory CpG
        oligonucleotides for treating and preventing infections in
        immunocompromised subjects)
IT
        (increasing immune response to; immunostimulatory CpG oligonucleotides
        for treating and preventing infections in immunocompromised subjects)
TΤ
    Dendritic cell
    Monocyte
        (induction of maturation of; immunostimulatory CpG oligonucleotides for
        treating and preventing infections in immunocompromised subjects)
IT
    Lentivirus
        (infection; immunostimulatory CpG oligonucleotides for treating and
        preventing infections in immunocompromised subjects)
IT
    Syphilis
        (neurosyphilis, opportunistic infection with; immunostimulatory CpG
        oligonucleotides for treating and preventing infections in
        immunocompromised subjects)
IT
    Lymphoma
        (non-Hodgkin's, systemic, opportunistic; immunostimulatory CpG
        oligonucleotides for treating and preventing infections in
        immunocompromised subjects)
TT
    Antigens
    RL: BSU (Biological study, unclassified); PAC (Pharmacological activity);
    THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (of immunodeficiency virus, in treatment of immunodeficiency virus
        infection; immunostimulatory CpG oligonucleotides for treating and
        preventing infections in immunocompromised subjects)
TТ
    Meningitis
        (opportunistic cryptococcal; immunostimulatory CpG oligonucleotides for
        treating and preventing infections in immunocompromised subjects)
TT
    Cryptosporidium
        (opportunistic cryptosporidiosis infection from; immunostimulatory CpG
        oligonucleotides for treating and preventing infections in
        immunocompromised subjects)
IT
        (opportunistic infection with Pneumocystis carinii; immunostimulatory
        CpG oligonucleotides for treating and preventing infections in
        immunocompromised subjects)
IT
    Cytomegalovirus
    Human herpesvirus
    Human herpesvirus 3
    Human papillomavirus
    Leishmania
    Molluscum contagiosum virus
    Mycobacterium
    Syphilis
        (opportunistic infection with; immunostimulatory CpG oligonucleotides
        for treating and preventing infections in immunocompromised subjects)
IT
    Hepatitis
    Histoplasma capsulatum
        (opportunistic infection; immunostimulatory CpG oligonucleotides for
        treating and preventing infections in immunocompromised subjects)
тт
    Isospora
        (opportunistic isosporiasis infection from; immunostimulatory CpG
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oligonucleotides for treating and preventing infections in
        immunocompromised subjects)
TТ
     Microsporidia
        (opportunistic microsporidiosis infection from; immunostimulatory CpG
        oligonucleotides for treating and preventing infections in
        immunocompromised subjects)
     Cryptococcus (fungus)
IT
        (opportunistic; immunostimulatory CpG oligonucleotides for treating and
        preventing infections in immunocompromised subjects)
IT
     Functional groups
        (phosphodiester; immunostimulatory CpG oligonucleotides for treating
        and preventing infections in immunocompromised subjects)
IT
     Pneumocystis carinii
        (pneumonia from; immunostimulatory CpG oligonucleotides for treating
        and preventing infections in immunocompromised subjects)
IT
     Brain, disease
        (progressive multifocal leukoencephalopathy, opportunistic;
        immunostimulatory CpG oligonucleotides for treating and preventing
        infections in immunocompromised subjects)
IT
     Infection
        (protozoal, as opportunistic infection; immunostimulatory CpG
        oligonucleotides for treating and preventing infections in
        immunocompromised subjects)
IT
     Mononuclear cell (leukocyte)
        (response to CpG oligodeoxynucleotides; immunostimulatory CpG
        oligonucleotides for treating and preventing infections in
        immunocompromised subjects)
     Salmonella
TT
        (salmonellosis from; immunostimulatory CpG oligonucleotides for
        treating and preventing infections in immunocompromised subjects)
IT
        (toxoplasmosis from, opportunistic infection; immunostimulatory CpG
        oligonucleotides for treating and preventing infections in
        immunocompromised subjects)
IT
     Neoplasm
     Prion diseases
        (treatment of; immunostimulatory CpG oligonucleotides for treating and
        preventing infections in immunocompromised subjects)
TT
     Antibodies and Immunoglobulins
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (vaccinated animals development of protective levels of;
        immunostimulatory CpG oligonucleotides for treating and preventing
        infections in immunocompromised subjects)
TT
     Immunization
        (vaccination; immunostimulatory CpG oligonucleotides for treating and
        preventing infections in immunocompromised subjects)
IT
     Infection
        (viral, as opportunistic infection; immunostimulatory CpG
        oligonucleotides for treating and preventing infections in
        immunocompromised subjects)
IT
     Interferons
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (.alpha., triggering secretion of; immunostimulatory CpG
        oligonucleotides for treating and preventing infections in
        immunocompromised subjects)
IT
     Interferons
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (.gamma., triggering secretion of; immunostimulatory CpG
        oligonucleotides for treating and preventing infections in
        immunocompromised subjects)
IT
     30516-87-1, AZT
     RL: BSU (Biological study, unclassified); PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (immunostimulatory CpG oligonucleotides for treating and preventing
        infections in immunocompromised subjects)
     2382-65-2
     RL: BSU (Biological study, unclassified); PAC (Pharmacological activity); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES
     (Uses)
        (immunostimulatory oligonucleotides containing; immunostimulatory CpG
        oligonucleotides for treating and preventing infections in
        immunocompromised subjects)
IT
     697319-80-5
                   697319-81-6
                                  697319-82-7 700878-94-0
     700878-95-1 700878-96-2
     RL: BSU (Biological study, unclassified); PAC (Pharmacological activity);
     PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES
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(Uses)
        (nucleotide sequence, immunocompromised macaques response to HBV
        vaccine improvement with; immunostimulatory CpG oligonucleotides for
        treating and preventing infections in immunocompromised subjects)
IT
     84216-59-1
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                               97802-24-9
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     RL: BSU (Biological study, unclassified); PAC (Pharmacological activity);
     PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES
     (Uses)
        (nucleotide sequence; immunostimulatory CpG oligonucleotides for
        treating and preventing infections in immunocompromised subjects)
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     698754-70-0 698754-71-1 698754-72-2
                                              698754-73-3
     698754-74-4
                   698754-75-5 698754-76-6 698754-77-7
     698754-85-7
                 698754-86-8 698754-87-9
    RL: PRP (Properties)
        (unclaimed nucleotide sequence; method of treating and preventing
        infections in immunocompromised subjects with immunostimulatory CpG
        oligonucleotides)
IT
    700878-94-0
    RL: BSU (Biological study, unclassified); PAC (Pharmacological activity);
     PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES
        (nucleotide sequence, immunocompromised macaques response to HBV
```

vaccine improvement with; immunostimulatory CpG oligonucleotides for

```
treating and preventing infections in immunocompromised subjects)
RN
     700878-94-0 HCAPLUS
     DNA, d(G-sp-G-sp-T-G-C-A-T-C-G-A-T-G-C-A-G-sp-G-sp-G-sp-G-sp-G-sp-G) (9CI)
CN
        (CA INDEX NAME)
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
     ANSWER 4 OF 21 HCAPLUS COPYRIGHT 2004 ACS on STN
L21
     2004:120676 HCAPLUS
DN
     140:175130
     Entered STN: 13 Feb 2004
ED
     Treating inflammatory arthropathies with oligodeoxynucleotide suppressors
TI
     of CpG-containing oligonucleotides
     Klinman, Dennis M.; Zeuner, Rainald; Verthelyi, Daniela
ΙN
     ; Gursel, Ihsan; Gursel, Mayda
PA
     The Government of the United States of America as Represented by the
     Secretary of the Department of Health and Human
     Services, USA
     PCT Int. Appl., 119 pp.
so
     CODEN: PIXXD2
DT
     Patent
LA
     English
IC
     ICM A61K
     1-7 (Pharmacology)
CC
FAN.CNT 2
                           KIND
                                  DATE
                                               APPLICATION NO.
                                                                        DATE
     PATENT NO.
     WO 2004012669
                           A2
                                  20040212
                                               WO 2003-US24205
                                                                        20030731
PΤ
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             GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
             LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM,
             PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
             KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR,
             BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
PRAI US 2002-400826P
                           Р
                                  20020801
     US 2002-401631P
                                  20020806
CLASS
 PATENT NO.
                  CLASS PATENT FAMILY CLASSIFICATION CODES
 WO 2004012669 'ICM
                         A61K
    The present disclosure relates to oligodeoxynucleotides that suppress an
     immune response. Methods are disclosed for preventing or treating
     inflammatory arthropathies by administering a therapeutically effective
     amount of a suppressive oligodeoxynucleotide. The oligodeoxynucleotides
     comprise .gtoreq.8 nucleotides in length, form a G-quadruplex (G tetrad), have a CD value of >2.9, has at least two guanosines, and suppresses an
     immune response. Optionally, the suppressive oligodeoxynucleotide has
     multiple guanosine-rich sequences, and in some examples, has one or more
     TTAGGG motifs; it may also be modified (e.g., with phosphorothioate
     linkages) to prevent degradation or is part of an oligodeoxynucleotide
     delivery complex that includes a targeting moiety. The suppressive
     oligodeoxynucleotides suppress CpG-DNA-induced immune activation, are
     highly specific, and are neither toxic nor non-specifically
     immunosuppressive. Suppression is mediated by the two-dimensional
     structure of the motifs, as shown by replacing individual guanosines by 7-deazaguanosine analogs to reduce G tetrad formation.
ST
     inflammatory arthropathy immunosuppression oligodeoxynucleotide; arthritis
     immunosuppression oligodeoxynucleotide
IT
     RL: ADV (Adverse effect, including toxicity); BIOL (Biological study)
         (CpG-containing, suppression of arthritis induced by; treating inflammatory
         arthropathies with oligodeoxynucleotide suppressors of CpG-containing
         oligonucleotides)
TT
     Steroids, biological studies
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (co-treatment with; treating inflammatory arthropathies with
         oligodeoxynucleotide suppressors of CpG-containing oligonucleotides)
     Joint, anatomical
IT
         (disease; treating inflammatory arthropathies with oligodeoxynucleotide
         suppressors of CpG-containing oligonucleotides)
ΤТ
     Drug delivery systems
         (liposomes; treating inflammatory arthropathies with
```

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oligodeoxynucleotide suppressors of CpG-containing oligonucleotides)
IT
     Anti-inflammatory agents
        (nonsteroidal, co-treatment with; treating inflammatory arthropathies
        with oligodeoxynucleotide suppressors of CpG-containing oligonucleotides)
     Collagens, biological studies
IT
     RL: ADV (Adverse effect, including toxicity); BIOL (Biological study)
        (suppression of arthritis induced by; treating inflammatory
        arthropathies with oligodeoxynucleotide suppressors of CpG-containing
        oligonucleotides)
    Lipids, biological studies RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (targeting moiety; treating inflammatory arthropathies with
        oligodeoxynucleotide suppressors of CpG-containing oligonucleotides)
     Anti-inflammatory agents
     Antiarthritics
     Antirheumatic agents
     Arthritis
     Human
     Immunosuppressants
     Rheumatoid arthritis
        (treating inflammatory arthropathies with oligodeoxynucleotide
        suppressors of CpG-containing oligonucleotides)
     Oligodeoxyribonucleotides
IT
     Phosphorothioate oligodeoxyribonucleotides
     RL: PAC (Pharmacological activity); PRP (Properties); THU (Therapeutic
     use); BIOL (Biological study); USES (Uses)
        (treating inflammatory arthropathies with oligodeoxynucleotide
        suppressors of CpG-containing oligonucleotides)
IT
    Drug delivery systems
        (virosomes, targeting moiety; treating inflammatory arthropathies with
        oligodeoxynucleotide suppressors of CpG-containing oligonucleotides)
IT
     329900-75-6, Cyclooxygenase 2
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (inhibitors, co-treatment with; treating inflammatory arthropathies
        with oligodeoxynucleotide suppressors of CpG-containing oligonucleotides)
     2382-65-2D, DNA containing
IT
     RL: ADV (Adverse effect, including toxicity); BIOL (Biological study)
        (suppression of arthritis induced by; treating inflammatory
        arthropathies with oligodeoxynucleotide suppressors of CpG-containing
        oligonucleotides)
     57-88-5, Cholesterol, biological studies
TT
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (targeting moiety; treating inflammatory arthropathies with
        oligodeoxynucleotide suppressors of CpG-containing oligonucleotides)
                                657710-64-0
IT
     157961-44-9
                  657710-63-9
                                              657710-65-1
                                                              657710-66-2
                   657710-68-4
                                 657710-69-5
                                               657710-70-8
     657710-67-3
                                                              657710-71-9
     657710-72-0
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                                               657710-75-3
                                                              657710-76-4
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                                 657710-79-7
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                   657710-83-3
                                 657710-84-4
                                               657710-85-5
                                                              657710-86-6
     657710-87-7
     RL: PAC (Pharmacological activity); PRP (Properties); THU (Therapeutic
     use); BIOL (Biological study); USES (Uses)
        (treating inflammatory arthropathies with oligodeoxynucleotide
        suppressors of CpG-containing oligonucleotides)
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     657711-37-0 657711-38-1 657711-39-2
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     657711-67-6 657711-68-7 657711-69-8
                              657711-72-3
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     657711-70-1 657711-71-2
     RL: PRP (Properties)
        (unclaimed nucleotide sequence; treating inflammatory arthropathies
        with oligodeoxynucleotide suppressors of CpG-containing oligonucleotides)
IT
     657710-97-9
     RL: PRP (Properties)
        (unclaimed nucleotide sequence; treating inflammatory arthropathies
        with oligodeoxynucleotide suppressors of CpG-containing oligonucleotides)
RN
     657710-97-9 HCAPLUS
CN
    DNA, d(N-N-N-R-Y-C-G-R-Y-N-N-G-G-G-G) (9CI) (CA INDEX NAME)
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
L21 ANSWER 5 OF 21 HCAPLUS COPYRIGHT 2004 ACS on STN
     2003:590824 HCAPLUS
AN
DN
    139:154891
     Entered STN: 01 Aug 2003
ED
    Multiple CpG oligodeoxynucleotides and their use to induce an immune
     response
TN
    Klinman, Dennis; Ishii, Ken; Verthelyi,
     Daniela
    The Government of the U.S.A., the Secretary of the Department of
    Health and Human Services, USA
    U.S. Pat. Appl. Publ., 41 pp., Cont.-in-part of Appl. No. PCT/US01/01122.
SO
     CODEN: USXXCO
DТ
     Patent
    English
LA
    ICM A61K048-00
IC
     ICS A61K031-70; C07H021-04
NCL
    514044000; 536023200
     63-6 (Pharmaceuticals)
FAN.CNT 2
     PATENT NO.
                         KIND DATE
                                            APPLICATION NO.
                                                                    DATE
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                                20030731
     US 2003144229
                         A1
                                            US 2002-194035
                                20010719
                                            WO 2001-US1122
     WO 2001051500
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         W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
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             HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT,
             LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN,
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         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
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                                20000114
PRAI US 2000-176115P
                          Р
    WO 2001-US1122
                          A2
                                20010112
CLASS
PATENT NO.
                 CLASS PATENT FAMILY CLASSIFICATION CODES
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US 2003144229 ICM
                        A61K048-00
                 ICS
                        A61K031-70; C07H021-04
                 NCL
                        514044000; 536023200
OS
    MARPAT 139:154891
    Compns. including multiple oligodeoxynucleotides with a CpG motif are disclosed herein. The compns. can include either D or K type
AB
    oligodeoxynucleotides. These compns. are of use in inducing an immune
     response in a large percentage of the individuals in a population.
ST
    CpG oligodeoxyribonucleotide sequence vaccine immunostimulant
    Antibodies and Immunoglobulins
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
    BIOL (Biological study); PREP (Preparation)
        (IgM, production of; multiple CpG oligodeoxynucleotides and their use to
        induce an immune response)
IT
     Immunostimulants
        (adjuvants; multiple CpG oligodeoxynucleotides and their use to induce
        an immune response)
IT
     Infection
        (bacterial; multiple CpG oligodeoxynucleotides and their use to induce
        an immune response)
IT
    Drug delivery systems
        (carriers; multiple CpG oligodeoxynucleotides and their use to induce
```

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an immune response)
IT
    Drug delivery systems
        (liposomes, targeting moiety; multiple CpG oligodeoxynucleotides and
        their use to induce an immune response)
IT
    Antibiotics
     Antiviral agents
     Fungicides
     Human
     Immunostimulants
     Immunotherapy
     Mycosis
     Neoplasm
     Vaccines
     Viral vectors
        (multiple CpG oligodeoxynucleotides and their use to induce an immune
        response)
TТ
     Oligodeoxyribonucleotides
     Phosphorothicate oligonucleotides
     RL: PAC (Pharmacological activity); PRP (Properties); THU (Therapeutic
     use); BIOL (Biological study); USES (Uses)
        (multiple CpG oligodeoxynucleotides and their use to induce an immune
        response)
IT
     Interleukin 6
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     BIOL (Biological study); PREP (Preparation)
        (production of; multiple CpG oligodeoxynucleotides and their use to induce
        an immune response)
ΙT
    Mononuclear cell (leukocyte)
        (proliferation of; multiple CpG oligodeoxynucleotides and their use to
        induce an immune response)
     Drug delivery systems
        (targeted; multiple CpG oligodeoxynucleotides and their use to induce
        an immune response)
IT
    Lipids, biological studies
     RL: PEP (Physical, engineering or chemical process); PYP (Physical
     process); THU (Therapeutic use); BIOL (Biological study); PROC (Process);
     USES (Uses)
        (targeting moiety; multiple CpG oligodeoxynucleotides and their use to
        induce an immune response)
IT
     Vaccines
        (tumor; multiple CpG oligodeoxynucleotides and their use to induce an
        immune response)
IT
    Antitumor agents
        (vaccines; multiple CpG oligodeoxynucleotides and their use to induce
        an immune response)
     Infection
TT
        (viral; multiple CpG oligodeoxynucleotides and their use to induce an
        immune response)
ΙT
     2382-65-2D, oligonucleotides
                                    569386-89-6
                                                  569386-90-9
                                                                 569386-91-0
     RL: PAC (Pharmacological activity); PRP (Properties); THU (Therapeutic
     use); BIOL (Biological study); USES (Uses)
        (multiple CpG oligodeoxynucleotides and their use to induce an immune
        response)
     57-88-5, Cholesterol, biological studies
     RL: PEP (Physical, engineering or chemical process); PYP (Physical
     process); THU (Therapeutic use); BIOL (Biological study); PROC (Process);
     USES (Uses)
        (targeting moiety; multiple CpG oligodeoxynucleotides and their use to
        induce an immune response)
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TΤ
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     569431-82-9
                    569431-83-0
                                   569431-84-1
                                                  569431-85-2
                                                                 569431-86-3
     569431-87-4
     RL: PRP (Properties)
         (unclaimed nucleotide sequence; multiple CpG oligodeoxynucleotides and
        their use to induce an immune response)
TT
     569431-00-1
     RL: PRP (Properties)
         (unclaimed nucleotide sequence; multiple CpG oligodeoxynucleotides and
        their use to induce an immune response)
RN
     569431-00-1 HCAPLUS
     DNA, d(G-G-T-G-C-A-C-C-G-A-T-G-C-A-G-G-G-G) (9CI) (CA INDEX NAME)
CN
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
L21 ANSWER 6 OF 21 HCAPLUS COPYRIGHT 2004 ACS on STN
AN
     2003:511473 HCAPLUS
     139:79147
ED
     Entered STN: 04 Jul 2003
     Use of CpG oligodeoxynucleotides to induce angiogenesis
TT
IN
     Klinman, Dennis M.; Zheng, Mei; Rouse, Barry T.
PΑ
     The Government of the United States of America as Represented by the
     Secretary of the Department of Health and Human
     Services, USA; The University of Tennessee Research Corporation PCT Int. Appl., 74 pp.
SO
     CODEN: PIXXD2
DΤ
     Patent
     English
LА
     ICM C12N
IC
     1-8 (Pharmacology)
CC
FAN.CNT 1
     PATENT NO.
                          KIND
                                 DATE
                                              APPLICATION NO.
                                                                       DATE
                                  -----
                                               -----
                          ----
                          A2
                                               WO 2002-US40955
PΙ
     WO 2003054161
                                 20030703
                                                                       20021219
     WO 2003054161
                           A3
                                  20031030
         W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
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             LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH,
             PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES,
              FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR, BF, BJ,
             CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
PRAI US 2001-343457P
                                  20011220
                           Р
CLASS
PATENT NO.
                 CLASS PATENT FAMILY CLASSIFICATION CODES
WO 2003054161
                 ICM
                         C12N
os
    MARPAT 139:79147
     This disclosure provides a method of inducing production of vascular
     endothelial growth factor by a cell. The method includes contacting the
     cell with a CpG oligonucleotide, thereby inducing the production of vascular
     endothelial growth factor by the cell. The disclosure further provides a
     method inducing neovascularization in a tissue. This method includes
     comprising introducing a CpG oligodeoxynucleotide into an area of the
     tissue wherein the formation of new blood vessels is desired, thereby
     inducing neovascularization in the area of the tissue.
ST
     CpG oligodeoxynucleotide angiogenesis induction
IT
     Immune system
        (cells; use of CpG oligodeoxynucleotides to induce angiogenesis by
        stimulating production of vascular endothelial growth factor in cells in
        relation to screening for antiangiogenic agents)
     Scalp
IT
     Transplant and Transplantation
        (neovascularization in; use of CpG oligodeoxynucleotides to induce
        angiogenesis by stimulating production of vascular endothelial growth
        factor in cells in relation to screening for antiangiogenic agents)
IT
     Angiogenesis
        (neovascularization; use of CpG oligodeoxynucleotides to induce
        angiogenesis by stimulating production of vascular endothelial growth
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factor in cells in relation to screening for antiangiogenic agents)
IT
    Human herpesvirus
    Human herpesvirus 1
        (oligodeoxynucleotides from; use of CpG oligodeoxynucleotides to induce
        angiogenesis by stimulating production of vascular endothelial growth
        factor in cells in relation to screening for antiangiogenic agents)
TΤ
    Blood vessel, disease
        (peripheral, treatment; use of CpG oligodeoxynucleotides to induce
        angiogenesis by stimulating production of vascular endothelial growth
        factor in cells in relation to screening for antiangiogenic agents)
TT
    Transplant and Transplantation
        (skin, neovascularization in; use of CpG oligodeoxynucleotides to
        induce angiogenesis by stimulating production of vascular endothelial
        growth factor in cells in relation to screening for antiangiogenic
        agents)
TT
     Skin
        (transplant, neovascularization in; use of CpG oligodeoxynucleotides to
        induce angiogenesis by stimulating production of vascular endothelial
        growth factor in cells in relation to screening for antiangiogenic
    Angiogenesis inhibitors
     Animal cell
     Blood vessel
     Drug screening
    Macrophage
        (use of CpG oligodeoxynucleotides to induce angiogenesis by stimulating
        production of vascular endothelial growth factor in cells in relation to
        screening for antiangiogenic agents)
    Oligodeoxyribonucleotides
    RL: BUU (Biological use, unclassified); PAC (Pharmacological activity);
    THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (use of CpG oligodeoxynucleotides to induce angiogenesis by stimulating
        production of vascular endothelial growth factor in cells in relation to
        screening for antiangiogenic agents)
TТ
    DNA
     Viral DNA
     RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (use of CpG oligodeoxynucleotides to induce angiogenesis by stimulating
        production of vascular endothelial growth factor in cells in relation to
        screening for antiangiogenic agents)
    Nucleic acids
     RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (viral; use of CpG oligodeoxynucleotides to induce angiogenesis by
        stimulating production of vascular endothelial growth factor in cells in
        relation to screening for antiangiogenic agents)
TT
    2382-65-2
     RL: BUU (Biological use, unclassified); PAC (Pharmacological activity);
     THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (oligodeoxynucleotides containing; use of CpG oligodeoxynucleotides to
        induce angiogenesis by stimulating production of vascular endothelial
        growth factor in cells in relation to screening for antiangiogenic
        agents)
IT
    552439-42-6
                  552439-43-7 552439-44-8
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     RL: PRP (Properties)
        (unclaimed nucleotide sequence; use of CpG oligodeoxynucleotides to
        induce angiogenesis)
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     RL: PRP (Properties)
        (unclaimed sequence; use of CpG oligodeoxynucleotides to induce
        angiogenesis)
     127464-60-2, Vascular endothelial growth factor
TT
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (use of CpG oligodeoxynucleotides to induce angiogenesis by stimulating
        production of vascular endothelial growth factor in cells in relation to
        screening for antiangiogenic agents)
TT
     550394-91-7
                    550394-92-8
     RL: BUU (Biological use, unclassified); PAC (Pharmacological activity);
     THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (use of CpG oligodeoxynucleotides to induce angiogenesis by stimulating
        production of vascular endothelial growth factor in cells in relation to
        screening for antiangiogenic agents)
IT
     552439-49-3
     RL: PRP (Properties)
        (unclaimed nucleotide sequence; use of CpG oligodeoxynucleotides to
        induce angiogenesis)
RN
     552439-49-3 HCAPLUS
    DNA, d(G-G-T-G-C-A-T-C-G-A-T-G-C-A-G-G-G-G-G) (9CI) (CA INDEX NAME)
CN
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
     ANSWER 7 OF 21 HCAPLUS COPYRIGHT 2004 ACS on STN
L21
     2003:377001 HCAPLUS
AN
DN
     138:390866
ED
     Entered STN: 16 May 2003
     Use of sterically stabilized cationic liposomes to efficiently deliver CpG
TT
     oligonucleotides in vivo
     Klinman, Dennis M.; Gursel, Ihsan; Ishii, Ken J.;
IN
     Kawakami, Koji; Joshi, Bharat H.; Puri, Raj K.
     Department of Health and Human Services, USA
PA
     PCT Int. Appl., 110 pp.
SO
     CODEN: PIXXD2
DT
     Patent
LA
     English
     ICM C12N
IC
     63-5 (Pharmaceuticals)
CC
     Section cross-reference(s): 8
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                                                                       DATE
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     WO 2003040308
                                  20030515
                                               WO 2002-US24235
PΤ
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                                                                       20020729
     WO 2003040308
                           A3
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             PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ,
             UA, UG, US, UZ, VN, YU, ZA, ZM, ZW
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
             KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF,
             CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
PRAI US 2001-308283P
                           P
                                  20010727
    US 2002-206407
                           Α
                                  20020725
CLASS
 PATENT NO.
                  CLASS PATENT FAMILY CLASSIFICATION CODES
WO 2003040308
                 ICM
                         C12N
    MARPAT 138:390866
     Sterically stabilized cationic liposomes (SSCL) encapsulating a K type
     oligodeoxynucleotide (ODN) including a CpG motif are disclosed. These
     SSCL encapsulating a K type ODN can be used to effectively deliver the ODN
     to a cell. A novel method is also disclosed for producing the SSCL encapsulating the K type ODN. Administration of the SSCL encapsulating a
     K type ODN and a chemotherapeutic agent, such as a chimeric mol.
     comprising a targeting mol. selected from the group consisting of an
     IL-13, and an anti-IL-13 receptor antibody; and an effector mol. selected
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from the group consisting of a Pseudomonas exotoxin, a Diphtheria toxin,
     and a radionuclide, can be used to dramatically reduce the growth of solid
     tumors.
     antitumor oligonucleotide CpG delivery liposome IL13 IL13R antibody
ST
     Infection
        (bacterial; use of sterically stabilized cationic liposomes to
        efficiently deliver CpG oligonucleotides in vivo)
IT
     Drug delivery systems
        (carriers; use of sterically stabilized cationic liposomes to
        efficiently deliver CpG oligonucleotides in vivo)
IT
     Toxins
     RL: PEP (Physical, engineering or chemical process); PYP (Physical
     process); THU (Therapeutic use); BIOL (Biological study); PROC (Process);
     USES (Uses)
        (diphtheria; use of sterically stabilized cationic liposomes to
        efficiently deliver CpG oligonucleotides in vivo)
TT
        (exotoxin; use of sterically stabilized cationic liposomes to
        efficiently deliver CpG oligonucleotides in vivo)
TT
     Toxins
     RL: PEP (Physical, engineering or chemical process); PYP (Physical
     process); THU (Therapeutic use); BIOL (Biological study); PROC (Process);
     USES (Uses)
        (exotoxins, of Pseudomonas; use of sterically stabilized cationic
        liposomes to efficiently deliver CpG oligonucleotides in vivo)
     Cytokines
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (induction of; use of sterically stabilized cationic liposomes to
        efficiently deliver CpG oligonucleotides in vivo)
     Drug delivery systems
        (injections, i.v.; use of sterically stabilized cationic liposomes to
        efficiently deliver CpG oligonucleotides in vivo)
TT
     Drug delivery systems
        (injections; use of sterically stabilized cationic liposomes to
        efficiently deliver CpG oligonucleotides in vivo)
IT
     Interleukin receptors
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (interleukin 13, antibodies to; use of sterically stabilized cationic
        liposomes to efficiently deliver CpG oligonucleotides in vivo)
IT
     Drug delivery systems
        (liposomes; use of sterically stabilized cationic liposomes to
        efficiently deliver CpG oligonucleotides in vivo)
IT
     Encapsulation
        (microencapsulation; use of sterically stabilized cationic liposomes to
        efficiently deliver CpG oligonucleotides in vivo)
IT
     Neck, anatomical
        (neoplasm; use of sterically stabilized cationic liposomes to
        efficiently deliver CpG oligonucleotides in vivo)
IT
    Neoplasm
        (solid; use of sterically stabilized cationic liposomes to efficiently
        deliver CpG oligonucleotides in vivo)
IT
     Drug delivery systems
        (targeted; use of sterically stabilized cationic liposomes to
        efficiently deliver CpG oligonucleotides in vivo)
     Antibodies and Immunoglobulins
IT
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (to IL13 receptor; use of sterically stabilized cationic liposomes to
        efficiently deliver CpG oligonucleotides in vivo)
IT
     Antimicrobial agents
     Antitumor agents
     Fungicides
     Head, neoplasm
     Human
     Immunostimulants
     Lymphocyte
     Mycosis
     Particle size distribution
     Stabilizing agents
     Vaccines
        (use of sterically stabilized cationic liposomes to efficiently deliver
        CpG oligonucleotides in vivo)
IT
     Interleukin 13
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (use of sterically stabilized cationic liposomes to efficiently deliver
        CpG oligonucleotides in vivo)
IT
     Polyoxyalkylenes, biological studies
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RL: MOA (Modifier or additive use); PEP (Physical, engineering or chemical
     process); POF (Polymer in formulation); PRP (Properties); PYP (Physical
     process); THU (Therapeutic use); BIOL (Biological study); PROC (Process);
     USES (Uses)
        (use of sterically stabilized cationic liposomes to efficiently deliver
        CpG oligonucleotides in vivo)
TT
    Aminoplasts
     Oligodeoxyribonucleotides
     Phosphatidylethanolamines, biological studies
     RL: PEP (Physical, engineering or chemical process); PRP (Properties); PYP
     (Physical process); THU (Therapeutic use); BIOL (Biological study); PROC
     (Process); USES (Uses)
        (use of sterically stabilized cationic liposomes to efficiently deliver
        CpG oligonucleotides in vivo)
TT
    Antigens
     Polysaccharides, biological studies
     Radionuclides, biological studies
    RL: PEP (Physical, engineering or chemical process); PYP (Physical
     process); THU (Therapeutic use); BIOL (Biological study); PROC (Process);
     USES (Uses)
        (use of sterically stabilized cationic liposomes to efficiently deliver
        CpG oligonucleotides in vivo)
TΤ
    Fusion proteins (chimeric proteins)
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (use of sterically stabilized cationic liposomes to efficiently deliver
        CpG oligonucleotides in vivo)
IT
    Infection
        (viral; use of sterically stabilized cationic liposomes to efficiently
        deliver CpG oligonucleotides in vivo)
    Interferons
    RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (.gamma., induction of; use of sterically stabilized cationic liposomes
        to efficiently deliver CpG oligonucleotides in vivo)
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     524986-14-9 524986-15-0 524986-16-1 524986-17-2
     RL: PRP (Properties)
        (unclaimed nucleotide sequence; use of sterically stabilized cationic
        liposomes to efficiently deliver CpG oligonucleotides in vivo)
TT
    25322-68-3, Polyethylene glycol
    RL: MOA (Modifier or additive use); PEP (Physical, engineering or chemical
    process); POF (Polymer in formulation); PRP (Properties); PYP (Physical
process); THU (Therapeutic use); BIOL (Biological study); PROC (Process);
    USES (Uses)
        (use of sterically stabilized cationic liposomes to efficiently deliver
        CpG oligonucleotides in vivo)
IT
    57-88-5, Cholesterol, biological studies 598-56-1
                                                           4004-05-1, Dioleoyl
     phosphatidyl ethanolamine
                                 9011-05-6, Carbamol
                                                      524974-46-7
    RL: PEP (Physical, engineering or chemical process); PRP (Properties); PYP
     (Physical process); THU (Therapeutic use); BIOL (Biological study); PROC
     (Process); USES (Uses)
        (use of sterically stabilized cationic liposomes to efficiently deliver
        CpG oligonucleotides in vivo)
IT
     2382-65-2D, oligonucleotides
    RL: PEP (Physical, engineering or chemical process); PYP (Physical
     process); THU (Therapeutic use); BIOL (Biological study); PROC (Process);
     USES (Uses)
        (use of sterically stabilized cationic liposomes to efficiently deliver
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CpG oligonucleotides in vivo)
ΙT
     524985-88-4
     RL: PRP (Properties)
         (unclaimed nucleotide sequence; use of sterically stabilized cationic
        liposomes to efficiently deliver CpG oligonucleotides in vivo)
RN
     524985-88-4 HCAPLUS
CN
     DNA, d(G-G-T-G-C-A-T-C-G-A-T-G-C-A-G-G-G-G-G) (9CI) (CA INDEX NAME)
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
L21 ANSWER 8 OF 21 HCAPLUS COPYRIGHT 2004 ACS on STN
     2003:306455 HCAPLUS
ΑN
DN
     139:159661
     Entered STN: 22 Apr 2003
     CpG Oligodeoxynucleotides Protect Normal and SIV-Infected Macaques from
     Leishmania Infection
ΑU
     Verthelyi, Daniela; Gursel, Mayda; Kenney, Richard T.;
     Lifson, Jeffrey D.; Liu, Shuying; Mican, Joan; Klinman, Dennis M.
     Center for Biologics Evaluation and Research, Section of Retroviral
CS
     Immunology, Food and Drug Administration, Bethesda, MD, 20892, USA Journal of Immunology (2003), 170(9), 4717-4723
SO
     CODEN: JOIMA3; ISSN: 0022-1767
     American Association of Immunologists
DT
     Journal
     English
LA
CC
     1-7 (Pharmacology)
     Oligodeoxynucleotides containing CpG motifs (CpG ODNs) mimic microbial DNA and
     activate effectors of the innate immune response, which limits the spread
     of pathogens and promotes an adaptive immune response. CpG ODNs have been
     shown to protect mice from infection with intracellular pathogens.
     Unfortunately, CpG motifs that optimally stimulate humans are only weakly
     active in mice, mandating the use of nonhuman primates to monitor the
     activity and safety of "human" CpG ODNs in vivo. This study demonstrates
     that CpG ODN treatment of rhesus macaques significantly reduces the
     severity of the lesions caused by a challenge with Leishmania: Leishmania
     superinfection is common in immunocompromised hosts, particularly those infected with HIV. This study shows that PBMCs from HIV-infected subjects
     respond to stimulation with CpG ODNs. To determine whether CpG ODNs can
     protect retrovirus-infected primates, SIV-infected macaques were treated
     with CpG ODNs and then challenged with Leishmania: Both lesion size and
     parasite load were significantly reduced in the CpG-treated animals. These findings support the clin. development of CpG ODNs as
     immunoprotective agents in normal and HIV-infected patients.
ST
     CpG oligodeoxynucleotide immunoprotection SIV Leishmania infection
     Human
     Immunostimulants
     Simian immunodeficiency virus
         (CpG oligodeoxynucleotides protect normal and SIV-infected macaques
         from Leishmania infection)
     Oligodeoxyribonucleotides
     Phosphorothioate oligodeoxyribonucleotides
     RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
         (CpG oligodeoxynucleotides protect normal and SIV-infected macaques
         from Leishmania infection)
TT
     Human immunodeficiency virus
         (CpG oligodeoxynucleotides protect normal and SIV-infected macaques
         from Leishmania infection in relation to immunoprotection in
        HIV-infected patients)
IT
     Leishmania
         (leishmaniasis from; CpG oligodeoxynucleotides protect normal and
        SIV-infected macaques from Leishmania infection)
     572925-91-8 572925-92-9
                                   572925-93-0 573722-80-2
     573722-82-4 573722-83-5
     RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
         (CpG oligodeoxynucleotides protect normal and SIV-infected macaques
        from Leishmania infection)
RE.CNT 46
               THERE ARE 46 CITED REFERENCES AVAILABLE FOR THIS RECORD
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(2) Azzoni, L; J Immunol 2002, V168, P5764 HCAPLUS
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    Southwestern Europe, 1990-1998 2000
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IT
     573722-80-2
     RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (CpG oligodeoxynucleotides protect normal and SIV-infected macaques
        from Leishmania infection)
RN
     573722-80-2 HCAPLUS
     DNA, d(G-sp-G-sp-T-G-C-A-T-C-G-A-T-G-C-A-G-sp-G-sp-G-sp-G-sp-G) (9CI)
CN
     (CA INDEX NAME)
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
L21 ANSWER 9 OF 21 HCAPLUS COPYRIGHT 2004 ACS on STN
     2003:241991 HCAPLUS
AN
DN
     138:270283
     Entered STN: 28 Mar 2003
     Oligodeoxynucleotide and its use to induce an immune response
IN
     Klinman, Dennis; Verthelyi, Daniela; Ishii,
     Ken; Mond, James J.; Gursel, Mayda
PΑ
     USA
     U.S. Pat. Appl. Publ., 52 pp., Cont.-in-part of U.S. Ser. No. 958,713.
SO
     CODEN: USXXCO
DT
     Patent
LA
     English
IC
     A61K048-00; C07H021-04
     514044000; 536023100
     15-2 (Immunochemistry)
     Section cross-reference(s): 1
FAN.CNT 2
     PATENT NO.
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                            A1
                                   20030327
                                                US 2002-68160
PT
     US 2003060440
                                                                          20020206
PRAI US 1999-128898P
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                                   19990412
     US 2001-958713
                                   20011011
CLASS
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CLASS PATENT FAMILY CLASSIFICATION CODES

PATENT NO.

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IC
                                          C07H021-04
 US 2003060440
                        A61K048-00IC
                 NCL
                        514044000; 536023100
                ECLA A61K031/7088; A61K031/7088+M; A61K039/39+M
 US 2003060440
     D type CpG oligodeoxynucleotides are provided herein that include a
     sequence represented by the following formula: 5'-X1X2X3PulPy2CpGPu3Py4X4X5X6(W)M(G)N-3' wherein the central CpG motif is
     unmethylated, Pu is a purine nucleotide, Py is a pyrimidine nucleotide, X
     and W are any nucleotide, M is any integer from 0 to 10, and N is any
     integer from 4 to 10. The oligodeoxynucleotides can activate immune
     cells, such as antigen-presenting cells or natural killer cell, and/or can
     stimulate production of cytokines. Methods of using these oligodeoxynucleotides to induce an immune response are provided.
     oligodeoxynucleotides can be used in treatment or amelioration of cancer,
     allergy, autoimmune disease, immunodeficiency, or infection. They can
     also be used to enhance the efficacy of vaccines.
ST
     CpG oligodeoxynucleotide immunotherapy immunostimulation
IT
     Allergy
     Allergy inhibitors
     Anti-infective agents
     Antibiotics
     Antitumor agents
     Antiviral agents
     Asthma
     Autoimmune disease
     Cell activation
     Chemotherapy
     Dendritic cell
     Fungicides
     Human
     Immunodeficiency
     Immunostimulation
     Immunotherapy
     Leishmania major
     Monocyte
     Mycosis
     Neoplasm
     Radiotherapy
     Vaccines
        (CpG oligodeoxynucleotides for immunostimulation and immunotherapy of
        various diseases)
IT
     Allergens
     RL: ADV (Adverse effect, including toxicity); BIOL (Biological study)
        (CpG oligodeoxynucleotides for immunostimulation and immunotherapy of
        various diseases)
TТ
     Cvtokines
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (CpG oligodeoxynucleotides for immunostimulation and immunotherapy of
        various diseases)
IT
     Interleukin 10
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (CpG oligodeoxynucleotides for immunostimulation and immunotherapy of
        various diseases)
     Oligodeoxyribonucleotides
     RL: BSU (Biological study, unclassified); PAC (Pharmacological activity);
     PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES
        (CpG-containing; CpG oligodeoxynucleotides for immunostimulation and
        immunotherapy of various diseases)
IT
     Immunostimulants
        (adjuvants; CpG oligodeoxynucleotides for immunostimulation and
        immunotherapy of various diseases)
IT
     Infection
        (bacterial; CpG oligodeoxynucleotides for immunostimulation and
        immunotherapy of various diseases)
IT
     Immunity
        (cell-mediated; CpG oligodeoxynucleotides for immunostimulation and
        immunotherapy of various diseases)
IT
     Lipids, biological studies
     RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (delivery complex; CpG oligodeoxynucleotides for immunostimulation and
        immunotherapy of various diseases)
IT
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (interferon .gamma.-inducible protein-10; CpG oligodeoxynucleotides for
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immunostimulation and immunotherapy of various diseases)
IT
    Drug delivery systems
        (liposomes; CpG oligodeoxynucleotides for immunostimulation and
        immunotherapy of various diseases)
IT
        (natural killer cell; CpG oligodeoxynucleotides for immunostimulation
        and immunotherapy of various diseases)
IT
    Neoplasm
        (solid; CpG oligodeoxynucleotides for immunostimulation and
        immunotherapy of various diseases)
IT
    Infection
        (viral; CpG oligodeoxynucleotides for immunostimulation and
        immunotherapy of various diseases)
IT
    Drug delivery systems
        (virosomes; CpG oligodeoxynucleotides for immunostimulation and
        immunotherapy of various diseases)
IT
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (.alpha.; CpG oligodeoxynucleotides for immunostimulation and
        immunotherapy of various diseases)
IT
    Interferons
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (.gamma.; CpG oligodeoxynucleotides for immunostimulation and
        immunotherapy of various diseases)
IT
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     503572-65-4 503572-66-5
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     PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES
        (CpG oligodeoxynucleotides for immunostimulation and immunotherapy of
        various diseases)
TT
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IT
     RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (delivery complex; CpG oligodeoxynucleotides for immunostimulation and
        immunotherapy of various diseases)
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IT
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        (unclaimed nucleotide sequence; oligodeoxynucleotide and its use to
        induce an immune response)
IT
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     RL: PRP (Properties)
        (unclaimed sequence; oligodeoxynucleotide and its use to induce an
        immune response)
IT
    503572-62-1
    RL: BSU (Biological study, unclassified); PAC (Pharmacological activity);
    PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES
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(Uses)
        (CpG oligodeoxynucleotides for immunostimulation and immunotherapy of
        various diseases)
     503572-62-1 HCAPLUS
RN
     DNA, d(G-G-T-G-C-A-T-C-G-A-T-G-C-A-G-G-G-G-G) (9CI) (CA INDEX NAME)
CN
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    ANSWER 10 OF 21 HCAPLUS COPYRIGHT 2004 ACS on STN
     2003:229621 HCAPLUS
AN
     139:20882
DN
ED
     Entered STN: 25 Mar 2003
TI
     Response of peripheral blood mononuclear cells from lupus patients to
     stimulation by CpG oligodeoxynucleotides
     Zeuner, R. A.; Klinman, D. M.; Illei, G.; Yarboro, C.; Ishii, K. J.; Gursel, M.; Verthelyi, D.
AII
CS
     Center for Biologics Evaluation and Research/Food and Drug Administration,
     Bethesda, MD, 20892, USA
     Rheumatology (Oxford, United Kingdom) (2003), 42(4), 563-569
SO
     CODEN: RUMAFK; ISSN: 1462-0324
PB
     Oxford University Press
DT
     Journal
     English
LA
     15-8 (Immunochemistry)
CC
AB
     Objectives: Increased levels of hypomethylated CpG-containing DNA in sera from
     patients with systemic lupus erythematosus (SLE) may contribute to the
     initiation and/or perpetuation of the disease. This study characterizes
     the in vitro response of peripheral blood mononuclear cells (PBMC) from SLE patients to CpG DNA. Methods: Secretion of cytokines and IgM, cell
     proliferation and up-regulation of co-stimulatory mols. were evaluated in
     PBMC from SLE patients and normal controls after stimulation with
     synthetic oligodeoxynucleotides (ODN) containing CpG motifs. Results:
     Up-regulation of co-stimulatory mols. and the secretion of interferon-.alpha. and interleukin-6 (IL-6) in response to CpG ODN was
     significantly reduced in monocytes and dendritic cells from SLE patients.
     Secretion of interferon-.gamma. by natural killer (NK) cells was also reduced. In contrast, the IgM and IL-10 response of B cells to CpG ODN
     was normal. Conclusion: Monocytes, dendritic cells and NK cells from SLE
     patients respond abnormally to CpG ODN stimulation, which may contribute
     to the cytokine imbalance observed in SLE.
     monocyte dendritic cell NK lupus CpG oligodeoxynucleotide cytokine
ST
TT
     Phosphorothioate oligodeoxyribonucleotides
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (CpG; monocytes, dendritic cells and NK cells from lupus patients
        respond to CpG oligodeoxynucleotides stimulation and contribute to the
        cytokine imbalance)
IT
     Histocompatibility antigens
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (HLA-DR; monocytes, dendritic cells and NK cells from lupus patients
        respond to CpG oligodeoxynucleotides stimulation and contribute to the
        cytokine imbalance)
IT
     Dendritic cell
     Monocyte
        (monocytes, dendritic cells and NK cells from lupus patients respond to
        CpG oligodeoxynucleotides stimulation and contribute to the cytokine
        imbalance)
IT
     CD40 (antigen)
     Interleukin 6
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (monocytes, dendritic cells and NK cells from lupus patients respond to
        CpG oligodeoxynucleotides stimulation and contribute to the cytokine
        imbalance)
IT
        (monocytes, dendritic cells and NK cells from lupus patients respond to
        CpG oligodeoxynucleotides stimulation which maycontribute to the
        cytokine imbalance)
IT
     Lymphocyte
        (natural killer cell; monocytes, dendritic cells and NK cells from
        lupus patients respond to CpG oligodeoxynucleotides stimulation and
        contribute to the cytokine imbalance)
IT
     Lupus erythematosus
        (systemic; monocytes, dendritic cells and NK cells from lupus patients
        respond to CpG oligodeoxynucleotides stimulation and contribute to the
        cytokine imbalance)
IT
     Interferons
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Search done by Noble Jarrell

RL: BSU (Biological study, unclassified); BIOL (Biological study)

(.alpha.; monocytes, dendritic cells and NK cells from lupus patients

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respond to CpG oligodeoxynucleotides stimulation and contribute to the
         cytokine imbalance)
IT
     Interferons
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
         (.qamma.; monocytes, dendritic cells and NK cells from lupus patients
         respond to CpG oligodeoxynucleotides stimulation and contribute to the
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RL: BSU (Biological study, unclassified); BIOL (Biological study)
         (monocytes, dendritic cells and NK cells from lupus patients respond to
         CpG oligodeoxynucleotides stimulation and contribute to the cytokine
         imbalance)
RE CNT
               THERE ARE 29 CITED REFERENCES AVAILABLE FOR THIS RECORD
        29
RE
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(28) Yabuhara, A; J Rheumatol 1996, V23, P171 MEDLINE (29) Yung, R; Lupus 1994, V3, P487 HCAPLUS
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     537058-58-5
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
         (monocytes, dendritic cells and NK cells from lupus patients respond to
         CpG oligodeoxynucleotides stimulation and contribute to the cytokine
         imbalance)
RN
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CN
     (CA INDEX NAME)
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
L21
     ANSWER 11 OF 21 HCAPLUS COPYRIGHT 2004 ACS on STN
     2003:202770 HCAPLUS
AN
     138:236936
DN
ED
     Entered STN: 14 Mar 2003
     Differentiation of human monocytes into mature functional dendritic cells
     with CpG oligodeoxynucleotides
     Klinman, Dennis M.; Gursel, Mayda; Verthelyi,
IN
     Daniela
PA
     The Government of the United States of America as Represented by the
     Secretary of Health and Human Services, USA
     PCT Int. Appl., 69 pp.
so
     CODEN: PIXXD2
DT
     Patent
LΑ
     English
IC
     ICM C12N
CC
     15-10 (Immunochemistry)
     Section cross-reference(s): 3, 9
FAN.CNT 1
     PATENT NO.
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WO 2003020884
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                                   20040916
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              LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH,
              PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD,
              RU, TJ, TM
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL,
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              NE, SN, TD, TG
     US 2004241841
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                                                                         20040212
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PRAI US 2001-312190P
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     WO 2002-US25732
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CLASS
 PATENT NO.
                  CLASS PATENT FAMILY CLASSIFICATION CODES
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                         ______
WO 2003020884 ICM
                          C12N
     This disclosure relates to dendritic cells, specifically to the methods of
     generating mature dendritic cells using D type oligodeoxynucleotides
     including a CpG motif. The methods include contacting a dendritic cell
     precursor with a D ODN to generate a mature dendritic cell. In one
     specific, non-limiting example, the method includes contacting the dendritic cell with an antigen. In another specific, non-limiting
     example, the method is a single step method wherein the D ODN is
     administered without other cytokines, such as GM-CSF and/or IL-4. These methods are of use both in vitro and in vivo. The findings disclosed
     herein document that mature DC can be rapidly and reproducibly generated
     by culturing PBMC or elutriated monocytes with D ODN in serum-free or
     conventional medium. These DC efficiently present antigen to autologous T
     cells in vitro and in vivo, and support the induction of Th1 biased immune
     responses. Without being bound by theory, the effect of D ODN on DC
     differentiation is dependent upon IFN.alpha. -secreting pDC being present
     during culture. Current results indicate that IFN.alpha. is necessary but not sufficient by itself to induce the differentiation of human monocytes
     into functionally active DC (Fig 2C). It is likely that functional
     differences between D and K type CpG ODN reflects differences in their
     recognition, uptake and/or processing by immune cells. These differences
     in uptake and activity have a structural basis. Whereas the
     immunostimulatory motif of a conventional ODN consists of a
     phosphorothicate TCGTT/A, the relevant motif in a D ODN consists of a
     phosphodiester purine/pyrimidine/CG/purine/pyrimidine hexamer (Verthelyi
     et al., Jlmmunol. 166: 2372-2377, 2001, which is incorporated herein by
     reference). In addition, the hexamer of a D ODN is flanked by complementary bases
     that form a hairpin loop with the CpG dinucleotide at its apex - secondary
     structure that is absent from conventional CpG ODN. Finally, D but not K
     ODN are capped at the 3' end with a poly-G tail. This poly-G tail may interact with scavenger receptors on immune cells. D ODN reproducibly
     stimulated approx. 60 % of monocytes to differentiate into DC, as determined
     phenotypically, histol. and functionally. The production of IFN.alpha. by pDC
     present in culture contributed to this maturation (Fig 1A and 2B). The DC
     generated by D ODN are functionally active, as they promote
     antigen-specific immune responses in vitro and in vivo (Fig 3, 4, 5 and
ST
     monocyte differentiation dendritic cell CpG oligonucleotide
     T cell (lymphocyte)
IT
         (activated, generating; differentiation of human monocytes into mature
         functional dendritic cells with CpG oligodeoxynucleotides)
IT
     Immunity
         (cell-mediated, producing; differentiation of human monocytes into
        mature functional dendritic cells with CpG oligodeoxynucleotides)
TT
     Ce11
         (contacting a dendritic cell precursor with; differentiation of human
        monocytes into mature functional dendritic cells with CpG
        oligodeoxynucleotides)
IT
     Antigens
     Cytokines
     Interleukin 4
     Peptides, biological studies
     Polysaccharides, biological studies
     Proteins
     RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
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(Uses)
        (contacting a dendritic cell precursor with; differentiation of human
        monocytes into mature functional dendritic cells with CpG
        oligodeoxynucleotides)
     Cell differentiation
     DNA sequences
     Dendritic cell
     Human
     Monocyte
        (differentiation of human monocytes into mature functional dendritic
        cells with CpG oligodeoxynucleotides)
TT
     Oligodeoxyribonucleotides
     Phosphorothioate oligodeoxyribonucleotides
     RL: BSU (Biological study, unclassified); BUU (Biological use,
     unclassified); PRP (Properties); BIOL (Biological study); USES (Uses)
        (differentiation of human monocytes into mature functional dendritic
        cells with CpG oligodeoxynucleotides)
     Oligonucleotides
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (dinucleotides, CpG; differentiation of human monocytes into mature
        functional dendritic cells with CpG oligodeoxynucleotides)
     Conformation
        (hairpin loop, oligodeoxynucleotides forming; differentiation of human
        monocytes into mature functional dendritic cells with CpG
        oligodeoxynucleotides)
IT
        (humoral, producing; differentiation of human monocytes into mature
        functional dendritic cells with CpG oligodeoxynucleotides)
IT
     Conformation
        (loop, nucleic acid, oligodeoxynucleotides forming; differentiation of
        human monocytes into mature functional dendritic cells with CpG
        oligodeoxynucleotides)
IT
     Immunity
        (producing; differentiation of human monocytes into mature functional
        dendritic cells with CpG oligodeoxynucleotides)
IT
     Interferons
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (.alpha., secretion of; differentiation of human monocytes into mature
        functional dendritic cells with CpG oligodeoxynucleotides)
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IT
     RL: BSU (Biological study, unclassified); BUU (Biological use,
     unclassified); PRP (Properties); BIOL (Biological study); USES (Uses)
        (ODN nucleotide sequence; differentiation of human monocytes into
        mature functional dendritic cells with CpG oligodeoxynucleotides)
TΤ
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     RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
        (contacting a dendritic cell precursor with; differentiation of human
        monocytes into mature functional dendritic cells with CpG
        oligodeoxynucleotides)
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     RL: PRP (Properties)
        (unclaimed nucleotide sequence; differentiation of human monocytes into
        mature functional dendritic cells with CpG oligodeoxynucleotides)
TT
    RL: BSU (Biological study, unclassified); BUU (Biological use,
    unclassified); PRP (Properties); BIOL (Biological study); USES (Uses) (ODN nucleotide sequence; differentiation of human monocytes into
        mature functional dendritic cells with CpG oligodeoxynucleotides)
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RN
CN
    DNA, d(G-G-T-G-C-A-T-C-G-A-T-G-C-A-G-G-G-G-G) (9CI) (CA INDEX NAME)
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
L21 ANSWER 12 OF 21 HCAPLUS COPYRIGHT 2004 ACS on STN
    2002:737476 HCAPLUS
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DN
     137:231323
     Entered STN: 29 Sep 2002
ED
     CpG oligodeoxynucleotides induce human monocytes to mature into functional
TI
     dendritic cells
AU
     Gursel, Mayda; Verthelyi, Daniela; Klinman,
     Dennis M.
     Section of Retroviral Immunology, Center for Biologics Evaluation and
CS
     Research, Food and Drug Administration, Bethesda, MD, 20892, USA
SO
     European Journal of Immunology (2002), 32(9), 2617-2622
     CODEN: EJIMAF; ISSN: 0014-2980
     Wiley-VCH Verlag GmbH & Co. KGaA
PB
DT
     Journal
LΑ
     English
CC
     15-10 (Immunochemistry)
     Dendritic cells (DC) excel at presenting antigen to T cells and thus make
ΔR
     a key contribution to the induction of primary and secondary immune
     responses. DC matured in vitro and pulsed with antigen show promise for
     the immunotherapy of cancer and infectious diseases. Synthetic oligonucleotides (ODN) expressing immunomodulatory "CpG motifs" were found
     to boost APC function in mice. Current results demonstrate that the
     recently identified "D" type of CpG ODN stimulate human peripheral blood
     monocytes to mature into functionally active DC over 2-4 days. The
     transition from monocyte to DC is characterized by the upregulation of
     CD83, CD86, CD80, CD40 and the down-regulation of CD14. These DC support
     antigen-specific humoral and cellular responses in vitro and in vivo. The
     differentiation of these monocytes is mediated by plasmacytoid DC, which respond to D type ODN by secreting IFN-.alpha.. Since D type CpG motifs
     are present in bacterial and viral DNA, the maturation of monocytes into
     functional DC may reflect a physiol. response that can be harnessed
     therapeutically through the use of CpG ODN.
     monocyte differentiation dendritic cell CpG oligonucleotide
ST
     Genetic element
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
         (CpG island; CpG oligodeoxynucleotides induce human monocytes to mature
         into functional dendritic cells)
     Antigen presentation
IT
     Cell differentiation
     Dendritic cell
     Human
     Monocyte
         (CpG oligodeoxynucleotides induce human monocytes to mature into
         functional dendritic cells)
     Oligodeoxyribonucleotides
     RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
     (Biological study)
         (CpG oligodeoxynucleotides induce human monocytes to mature into
         functional dendritic cells)
ΤT
     T cell (lymphocyte)
         (activation; CpG oligodeoxynucleotides induce human monocytes to mature
         into functional dendritic cells)
IT
     Interferons
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
         (.alpha.; CpG oligodeoxynucleotides induce human monocytes to mature
         into functional dendritic cells)
     459471-16-0 459471-17-1 459471-18-2
     RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
     (Biological study)
         (CpG oligodeoxynucleotides induce human monocytes to mature into
         functional dendritic cells)
RE.CNT
               THERE ARE 36 CITED REFERENCES AVAILABLE FOR THIS RECORD
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IT
     Interleukin 6
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
         (differential and competitive activation of human immune cells by
         distinct classes of CpG oligodeoxynucleotide and formation of)
IT
     Monocyte
         (differentiation; differential and competitive activation of human
         immune cells by distinct classes of CpG oligodeoxynucleotide)
IT
     Lymphocyte
         (natural killer cell; differential and competitive activation of human
         immune cells by distinct classes of CpG oligodeoxynucleotide)
     Interferons
IT
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
         (.gamma.; differential and competitive activation of human immune cells
         by distinct classes of CpG oligodeoxynucleotide and formation of)
IT
     434529-76-7 434529-77-8
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
         (differential and competitive activation of human immune cells by
         distinct classes of CpG oligodeoxynucleotide)
RE.CNT
               THERE ARE 40 CITED REFERENCES AVAILABLE FOR THIS RECORD
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     434529-77-8
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
         (differential and competitive activation of human immune cells by
         distinct classes of CpG oligodeoxynucleotide)
RN
     434529-77-8 HCAPLUS
     DNA, d(G-sp-G-T-G-C-A-T-C-G-A-T-G-C-A-G-G-G-G-Sp-G) (9CI) (CA INDEX
CN
     NAME)
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
     ANSWER 14 OF 21 HCAPLUS COPYRIGHT 2004 ACS on STN
L21
     2002:124395 HCAPLUS
AN
DN
     136:293135
ED
     Entered STN: 17 Feb 2002
     CpG oligodeoxynucleotides as vaccine adjuvants in primates
TT
     Verthelyi, Daniela; Kenney, Richard T.; Seder, Robert A.; Gam,
ΑU
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Albert A.; Friedag, Brenda; Klinman, Dennis M.
     Division of Viral Products, Center for Biologics Evaluation and
CS
     Research/Food and Drug Administration, Bethesda, MD, 20892, USA
SO
     Journal of Immunology (2002), 168(4), 1659-1663
     CODEN: JOIMA3; ISSN: 0022-1767
PB
     American Association of Immunologists
DT
     Journal
     English
LΑ
CC
     15-2 (Immunochemistry)
     Synthetic oligodeoxynucleotides (ODN) containing unmethylated CpG motifs act
AB
     as immune adjuvants in mice, boosting the humoral and cellular response to
     coadministered Ags. CpG ODN that stimulate human PBMC are only weakly
     active in mice. Thus, alternative animal models are needed to monitor the activity and safety of "human" CpG ODN in vivo. This work demonstrates
     that rhesus macaques recognize and respond to the same CpG motifs that
     trigger human immune cells. Coadministering CpG ODN with heat-killed
     Leishmania vaccine provided significantly increased protection of macaques
     against cutaneous Leishmania infection. These findings indicate that
     rhesus macaques provide a useful model for studying the in vivo activity
     of human CpG motifs, and that ODN expressing these motifs act as strong
     immune adjuvants.
ST
     vaccine adjuvant CpG oligodeoxynucleotide Macaca Leishmania
TT
     Human
     Leishmania major
     Macaca mulatta
     Vaccines
         (CpG oligodeoxynucleotides as vaccine adjuvants in primates)
IT
     Interleukin 6
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
         (CpG oligodeoxynucleotides as vaccine adjuvants in primates)
     Oligodeoxyribonucleotides
IT
     RL: PAC (Pharmacological activity); PRP (Properties); THU (Therapeutic
     use); BIOL (Biological study); USES (Uses)
         (CpG oligodeoxynucleotides as vaccine adjuvants in primates)
     Antibodies and Immunoglobulins
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
         (IgG; CpG oligodeoxynucleotides as vaccine adjuvants in primates)
     Immunostimulants
IT
         (adjuvants; CpG oligodeoxynucleotides as vaccine adjuvants in primates)
IT
     T cell (lymphocyte)
         (proliferation; CpG oligodeoxynucleotides as vaccine adjuvants in
        primates)
IT
     Interferons
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
         (.gamma.; CpG oligodeoxynucleotides as vaccine adjuvants in primates)
IT
     406966-34-5 406966-35-6 406966-36-7 406966-37-8
     RL: PAC (Pharmacological activity); PRP (Properties); THU (Therapeutic
     use); BIOL (Biological study); USES (Uses)
        (CpG oligodeoxynucleotides as vaccine adjuvants in primates)
RE.CNT
        38
               THERE ARE 38 CITED REFERENCES AVAILABLE FOR THIS RECORD
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    406966-34-5
     RL: PAC (Pharmacological activity); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (CpG oligodeoxynucleotides as vaccine adjuvants in primates)
RN
     406966-34-5 HCAPLUS
      DNA, \ d(G-sp-G-sp-T-G-C-A-C-C-G-G-T-G-C-A-G-G-sp-G-sp-G-sp-G-sp-G) \ (9C1) 
CN
     (CA INDEX NAME)
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
    ANSWER 15 OF 21 HCAPLUS COPYRIGHT 2004 ACS on STN
L21
     2002:122818 HCAPLUS
AN
DN
     136:182447
ED
     Entered STN: 15 Feb 2002
     Vaccine against respiratory syncytial virus (RSV)
TT
IN
     Mond, James J.; Prince, Gregory; Klinman, Dennis M.
     Henry M. Jackson Foundation for the Advancement of Military Medicine, USA
so
     PCT Int. Appl., 30 pp.
     CODEN: PIXXD2
DT
     Patent
LΑ
     English
IC
     ICM A61K039-39
     15-2 (Immunochemistry)
CC
     Section cross-reference(s): 3, 63
FAN.CNT 1
     PATENT NO.
                          KIND
                                 DATE
                                              APPLICATION NO.
                                                                       DATE
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                          ----
    WO 2002011761
                          A2
                                  20020214
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PΤ
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         W: AU, CA, JP, US
         RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR
                           A5
                                  20020218
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PRAI US 2000-224011P
                           Р
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     US 2000-229307P
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     WO 2001-US41633
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CLASS
 PATENT NO.
                 CLASS PATENT FAMILY CLASSIFICATION CODES
 WO 2002011761 ICM
                         A61K039-39
     The present invention relates to a vaccine comprising adjuvanting
     oligodideoxynucleotides (ODNs), containing at least one CpG dinucleotide and
     an antigen comprising a peptide sequence bearing at least one epitope of a
     Paramyxoviridae F protein. In one embodiment, the ODN is admixed or
     conjugated to an F protein from a respiratory syncytial virus (RSV).
     vaccine of the invention may be administered directly to mucosal tissues
     of the respiratory tract by inhalation or intranasal administration.
ST
     vaccine Pneumovirus Paramyxoviridae F protein CpG; immunostimulatory
     oligonucleotide respiratory syncytial virus vaccine
IT
     Oligonucleotides
     RL: BSU (Biological study, unclassified); PRP (Properties); THU
     (Therapeutic use); BIOL (Biological study); USES (Uses)
        (CpG-containing immunostimulatory; vaccine compns. comprising
        Paramyxoviridae F protein epitopes and immunostimulatory
        oligonucleotides against respiratory syncytial virus)
IT
     Proteins
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES
     (Uses)
        (F; vaccine compns. comprising Paramyxoviridae F protein epitopes and
        immunostimulatory oligonucleotides against respiratory syncytial virus)
IT
     Immunostimulants
        (adjuvants; vaccine compns. comprising Paramyxoviridae F protein
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epitopes and immunostimulatory oligonucleotides against respiratory
        syncytial virus)
IT
    Respiratory tract
        (administration; vaccine compns. comprising Paramyxoviridae f protein
        epitopes and immunostimulatory oligonucleotides against respiratory
        syncytial virus)
TT
     Proteins
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES
     (Uses)
        (conjugates, G/F proteins; vaccine compns. comprising Paramyxoviridae F
        protein epitopes and immunostimulatory oligonucleotides against
        respiratory syncytial virus)
IT
    Drug delivery systems
        (inhalants; vaccine compns. comprising Paramyxoviridae F protein
        epitopes and immunostimulatory oligonucleotides against respiratory
        syncytial virus)
    Drug delivery systems
IT
        (injections, i.m.; vaccine compns. comprising Paramyxoviridae F protein
        epitopes and immunostimulatory oligonucleotides against respiratory
        syncytial virus)
IT
    Drug delivery systems
        (mucosal; vaccine compns. comprising Paramyxoviridae F protein epitopes
        and immunostimulatory oligonucleotides against respiratory syncytial
        virus)
IT
    Drug delivery systems
        (nasal, intra-; vaccine compns. comprising Paramyxoviridae F protein
        epitopes and immunostimulatory oligonucleotides against respiratory
        syncytial virus)
IT
     Drug delivery systems
        (respiratory tract; vaccine compns. comprising Paramyxoviridae F
        protein epitopes and immunostimulatory oligonucleotides against
        respiratory syncytial virus)
IT
    Drug delivery systems
        (solns., nasal; vaccine compns. comprising Paramyxoviridae F protein
        epitopes and immunostimulatory oligonucleotides against respiratory
        syncytial virus)
     DNA sequences
ΙT
     Epitopes
     Paramyxoviridae
     Pneumovirus
     Respiratory syncytial virus
     Vaccines
        (vaccine compns. comprising Paramyxoviridae F protein epitopes and
        immunostimulatory oligonucleotides against respiratory syncytial virus)
    Fusion proteins (chimeric proteins)
     G proteins (guanine nucleotide-binding proteins)
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES
     (Uses)
        (vaccine compns. comprising Paramyxoviridae F protein epitopes and
        immunostimulatory oligonucleotides against respiratory syncytial virus)
IT
    Antigens
     RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (vaccine compns. comprising Paramyxoviridae F protein epitopes and
        immunostimulatory oligonucleotides against respiratory syncytial virus)
     2382-65-2D, oligonucleotide derivs. 185810-20-2 207496-24-0
IT
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     Immunostimulatory RNA/DNA hybrid molecules
     Mond, James J.; Flora, Michael; Klinman, Dennis M.
IN
PΑ
     Biosynexus Incorporated, USA
SO
     PCT Int. Appl., 68 pp.
     CODEN: PIXXD2
DT
     Patent
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     English
     ICM A61K039-00
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     15-2 (Immunochemistry)
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                         4B024/GA11; 4B024/HA17; 4C084/AA13; 4C084/BA35;
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4C085/FF30; 4C086/AA01; 4C086/AA02; 4C086/EA16;
                         4C086/MA03; 4C086/MA04; 4C086/NA14; 4C086/ZB09
     The present invention provides immunol. compns. and methods relating to
     immunostimulatory intra-strand DNA/RNA hybrid oligonucleotides (HDRs),
     optimally encoding one or more CpG motif, which may be an unmethylated CpG
     motif. Administration of these compds., alone or in the context of one or
     more target antigens, promotes innate and antigen specific immunities.
ST
     vaccine adjuvant immunostimulatory RNA DNA hybrid antigen cytokine Ig
IT
     Genetic element
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (CpG island; immunostimulatory RNA/DNA hybrid mols. for use in
        vaccines)
IT
     Oligonucleotides
     RL: PAC (Pharmacological activity); PRP (Properties); THU (Therapeutic
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use); BIOL (Biological study); USES (Uses)
        (CpG-containing; immunostimulatory RNA/DNA hybrid mols. for use in
        vaccines)
IT
     Recombination, genetic
        (Ig class switching; immunostimulatory RNA/DNA hybrid mols. for use in
        vaccines)
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    Antibodies and Immunoglobulins
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (IqA; immunostimulatory RNA/DNA hybrid mols. for use in vaccines)
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        (IgG1; immunostimulatory RNA/DNA hybrid mols. for use in vaccines)
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        (IgG2; immunostimulatory RNA/DNA hybrid mols. for use in vaccines)
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TT
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        (IgM; immunostimulatory RNA/DNA hybrid mols. for use in vaccines)
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     Immunostimulants
        (adjuvants; immunostimulatory RNA/DNA hybrid mols. for use in vaccines)
IΤ
     Immunity
        (cell-mediated; immunostimulatory RNA/DNA hybrid mols. for use in
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IT
    T cell (lymphocyte)
        (helper cell/inducer, TH1; immunostimulatory RNA/DNA hybrid mols. for
        use in vaccines)
IT
     T cell (lymphocyte)
        (helper cell/inducer, TH2; immunostimulatory RNA/DNA hybrid mols. for
        use in vaccines)
TΨ
    Immunity
        (humoral; immunostimulatory RNA/DNA hybrid mols. for use in vaccines)
TT
     Immunostimulants
     Vaccines
        (immunostimulatory RNA/DNA hybrid mols. for use in vaccines)
    Antibodies and Immunoglobulins
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     Interleukin 12
     Interleukin 6
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     RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (immunostimulatory RNA/DNA hybrid mols. for use in vaccines)
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     Lymphocyte
        (natural killer cell; immunostimulatory RNA/DNA hybrid mols. for use in
        vaccines)
IT
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     T cell (lymphocyte)
        (proliferation; immunostimulatory RNA/DNA hybrid mols. for use in
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        (.gamma.; immunostimulatory RNA/DNA hybrid mols. for use in vaccines)
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     RL: PAC (Pharmacological activity); PRP (Properties); THU (Therapeutic
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IT
     381260-71-5
    RL: PRP (Properties)
        (unclaimed nucleotide sequence; immunostimulatory RNA/DNA hybrid mols.)
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    RNA, (G-G-U-G-C-A-U-C-G-A-U-G-C-A-G-G-G-G) (9CI) (CA INDEX NAME)
CN
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
    ANSWER 17 OF 21 HCAPLUS COPYRIGHT 2004 ACS on STN
L21
    2001:526086 HCAPLUS
DN
    135:102560
    Entered STN: 20 Jul 2001
ED
    Oligodeoxynucleotide and its use to induce an immune response
ΤI
IN
    Klinman, Dennis; Ishii, Ken; Verthelyi,
    Daniela
PA
    United States Dept. of Health and Human
    Services, USA
SO
    PCT Int. Appl., 48 pp.
     CODEN: PIXXD2
DT
    Patent
    English
T.A
    ICM C07H021-00
IC
     ICS A61K031-70; A61K039-39; C12N015-11
    1-7 (Pharmacology)
    Section cross-reference(s): 15, 63
FAN. CNT 2
    PATENT NO.
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                                DATE
                                            APPLICATION NO.
                                                                    DATE
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    WO 2001051500
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                                            WO 2001-US1122
PΙ
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             LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU,
             SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
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                                              AU 2001-27889
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                                  20010724
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     EP 1322655
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                                  20030702
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     US 2003144229
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                           A1
                                                                       20020712
PRAI US 2000-176115P
                                  20000114
                                  20010112
     WO 2001-US1122
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CLASS
 PATENT NO.
                  CLASS PATENT FAMILY CLASSIFICATION CODES
                  ICM
                         C07H021-00
 WO 2001051500
                         A61K031-70; A61K039-39; C12N015-11
                  ICS
     The present invention provides a substantially pure or isolated
     oligodeoxynucleotide (ODN) of at least about 10 nucleotides comprising
     different CpG sequences, as well as an oligodeoxynucleotide delivery
     complex and a pharmacol. composition comprising an ODN or ODNs, and a method of
     inducing an immune response by administering such an ODN or ODNs to a
     host.
     phosphorothioate oligodeoxynucleotide immunostimulant antitumor sequence
ST
IT
     Genetic element
     RL: PEP (Physical, engineering or chemical process); THU (Therapeutic
     use); BIOL (Biological study); PROC (Process); USES (Uses)
         (CpG island; oligodeoxynucleotide and its use to induce an immune
        response)
IT
     B cell (lymphocyte)
         (activation; oligodeoxynucleotide and its use to induce an immune
        response)
     Immunostimulants
IT
         (adjuvants; oligodeoxynucleotide and its use to induce an immune
        response)
     Chemical warfare agents
IT
         (biowarfare agents; oligodeoxynucleotide and its use to induce an
        immune response)
IT
     Immunity
        (cell-mediated; oligodeoxynucleotide and its use to induce an immune
        response)
IT
     Lipids, biological studies
     RL: PEP (Physical, engineering or chemical process); THU (Therapeutic
     use); BIOL (Biological study); PROC (Process); USES (Uses)
         (drug targeting with; oligodeoxynucleotide and its use to induce an
        immune response)
IT
     Immunity
        (humoral; oligodeoxynucleotide and its use to induce an immune
        response)
     Drug delivery systems
IT
         (liposomes, drug targeting with; oligodeoxynucleotide and its use to
        induce an immune response)
IT
     Allergy inhibitors
     Antiasthmatics
     Antibiotics
     Antitumor agents
     Drug targeting
     Immunodeficiency
     Immunostimulants
     Vaccines
        (oligodeoxynucleotide and its use to induce an immune response)
     Oligodeoxyribonucleotides
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); PEP (Physical, engineering or chemical process); PRP
     (Properties); THU (Therapeutic use); BIOL (Biological study); PROC
     (Process); USES (Uses)
         (oligodeoxynucleotide and its use to induce an immune response)
TΤ
     Antibodies
     Cytokines
     Interleukin 6
     RL: BSU (Biological study, unclassified); MFM (Metabolic formation); BIOL
     (Biological study); FORM (Formation, nonpreparative)
         (oligodeoxynucleotide and its use to induce an immune response)
     Phosphorothioate oligodeoxyribonucleotides
     RL: PEP (Physical, engineering or chemical process); THU (Therapeutic use); BIOL (Biological study); PROC (Process); USES (Uses)
        (oligodeoxynucleotide and its use to induce an immune response)
IT
     Neoplasm
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(solid, inhibitors; oligodeoxynucleotide and its use to induce an
        immune response)
IT
    Drug delivery systems
        (virosomes, drug targeting with; oligodeoxynucleotide and its use to
        induce an immune response)
    Interferons
TT
     RL: BSU (Biological study, unclassified); MFM (Metabolic formation); BIOL
     (Biological study); FORM (Formation, nonpreparative)
        (.gamma.; oligodeoxynucleotide and its use to induce an immune
        response)
IT
    57-88-5, Cholesterol, biological studies
     RL: PEP (Physical, engineering or chemical process); THU (Therapeutic
    use); BIOL (Biological study); PROC (Process); USES (Uses)
        (drug targeting with; oligodeoxynucleotide and its use to induce an
        immune response)
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     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (oligodeoxynucleotide and its use to induce an immune response)
IT
    2382-65-2
    RL: PEP (Physical, engineering or chemical process); THU (Therapeutic use); BIOL (Biological study); PROC (Process); USES (Uses)
        (oligodeoxynucleotide and its use to induce an immune response)
              THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT
(1) Cpg Immunopharmaceuticals Gmbh; WO 9958118 A 1999 HCAPLUS
(2) Davis, H; WO 9840100 A 1998 HCAPLUS
(3) Dynavax Technologies Corp; WO 9855495 A 1998 HCAPLUS
(4) Hybridon Inc; WO 9849288 A 1998 HCAPLUS
(5) Ottawa Civic Loeb Res Inst; WO 9956755 A 1999 HCAPLUS
(6) Univ Iowa Res Found; WO 9818810 A 1998 HCAPLUS
(7) Univ Iowa Res Found; WO 9951259 A 1999 HCAPLUS
(8) Wu, T; WO 9852581 A 1998 HCAPLUS
    350270-92-7
    RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (oligodeoxynucleotide and its use to induce an immune response)
RΝ
     350270-92-7 HCAPLUS
CN
    DNA, d(G-sp-G-T-G-C-A-C-C-G-A-T-G-C-A-G-G-G-sp-G) (9CI) (CA INDEX NAME)
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
    ANSWER 18 OF 21 HCAPLUS COPYRIGHT 2004 ACS on STN
L21
     2001:129322 HCAPLUS
AN
DN
    135:302807
                  21 Feb 2001
ED
    Entered STN:
ΤI
    Response of porcine peripheral blood mononuclear cells to CpG-containing
    oligodeoxynucleotides
ΑU
    Kamstrup, S.; Verthelyi, D.; Klinman, D. M.
    Department for Pathobiology and Diagnostics, Danish Veterinary Institute
CS
     for Virus Research, Kalvehave, Lindholm, DK-4771, Den.
    Veterinary Microbiology (2001), 78(4), 353-362
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CODEN: VMICDQ; ISSN: 0378-1135
     Elsevier Science B.V.
PB
DT
     Journal
LΑ
     English
CC
     15-10 (Immunochemistry)
     Exposure to bacterial DNA generates a "danger signal" that stimulates
     cellular elements of the mammalian immune system to proliferate and/or
     secrete cytokines. Stimulation is critically dependent on hexameric
     motifs that contain an unmethylated CpG dinucleotide: these are commonly
     found in bacterial but not vertebrate DNA. Different motifs are optimally stimulatory in different species. This work examines whether
     oligodeoxynucleotides (ODNs) containing CpG motifs stimulate peripheral blood
     mononuclear cells from pigs. Results show that pigs respond to CpG ODN by
     proliferating and secreting IL-6, IL-12 and TNF-.alpha.. By screening a
     large panel (>100) of ODNs, the palindromic hexamer 'ATCGAT' was
     identified as being optimally active in all animals examined (N=10).
      findings are the first to establish the immunostimulatory activity of CpG
     ODN in pigs, and suggest that the therapeutic uses envisioned for these
     ODNs (as vaccine adjuvants and immunoprotective agents) may be applicable
     to husbandry animals.
     CpG oligodeoxynucleotide adjuvant leukocyte proliferation cytokine swine
TT
     Genetic element
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
         (CpG island; response of porcine peripheral blood mononuclear cells to
         CpG-containing oligodeoxynucleotides)
·IT
     Oligodeoxyribonucleotides
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
      study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL
      (Biological study); USES (Uses)
       : (CpG-containing; response of porcine peripheral blood mononuclear cells to
         CpG-containing oligodeoxynucleotides)
TΤ
     Immunostimulants
         (adjuvants; response of porcine peripheral blood mononuclear cells to
         CpG-containing oligodeoxynucleotides)
     Cell proliferation
     Mononuclear cell (leukocyte)
     Swine
     Vaccines
         (response of porcine peripheral blood mononuclear cells to CpG-containing
         oligodeoxynucleotides)
     Interleukin 12
      Interleukin 6
     Tumor necrosis factors
     RL: BSU (Biological study, unclassified); MFM (Metabolic formation); BIOL (Biological study); FORM (Formation, nonpreparative)
         (response of porcine peripheral blood mononuclear cells to CpG-containing
         oligodeoxynucleotides)
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); PRP (Properties); BIOL (Biological study)
         (CpG oligodeoxynucleotides containing; response of porcine peripheral blood
         mononuclear cells to CpG-containing oligodeoxynucleotides)
IT
     366065-16-9
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL
      (Biological study); USES (Uses)
         (response of porcine peripheral blood mononuclear cells to CpG-containing oligodeoxynucleotides)
RE.CNT
               THERE ARE 32 CITED REFERENCES AVAILABLE FOR THIS RECORD
         32
RE
(1) Ballas, Z; J Immunol 1996, V157, P1840 HCAPLUS
(2) Bauer, M; Immunology 1999, V97(4), P699 HCAPLUS(3) Brown, W; Infect Immunol 1998, V66(11), P5423 HCAPLUS
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(17) Krieg, A; Antisense Nucleic Acid Drug Dev 1996, V6, P133 HCAPLUS (18) Krieg, A; Nature 1995, V374(6522), P546 HCAPLUS (19) Liang, H; J Clin Invest 1996, V98(5), P1119 HCAPLUS (20) McCluskie, M; J Immunol 1998, V161(9), P4463 HCAPLUS (21) McCluskie, M; Vaccine 1999, V18(3-4), P231 HCAPLUS (22) Moss, R; Vaccine 2000, V18(11-12), P1081 MEDLINE (23) Oxenius, A; J Virol 1999, V73(5), P4120 HCAPLUS (24) Pisetsky, D; Immunopharmacology 1998, V40(3), P199 HCAPLUS (25) Stacey, K; Infect Immun 1999, V678, P3719 (26) Walker, P; Proc Natl Acad Sci USA 1999, V96(12), P6970 HCAPLUS (27) Weiner, G; Proc Natl Acad Sci USA 1997, V9420, P10833 (28) Wloch, M; Human Gene Therapy 1998, V9, P1439 MEDLINE (29) Yamamoto, T; Jpn J Cancer Res 1994, V85, P775 HCAPLUS (30) Yi, A; J Immunol 1996, V157(12), P5394 HCAPLUS (31) Yi, A; J Immunol 1998, V161(9), P4493 HCAPLUS (32) Zimmermann, S; J Immunol 1998, V1608, P3627 366065-16-9 RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (response of porcine peripheral blood mononuclear cells to CpG-containing oligodeoxynucleotides) 366065-16-9 HCAPLUS RN $\label{eq:defDNA} DNA, \ d\left(G-sp-G-sp-T-G-C-A-T-C-G-A-T-G-C-A-G-sp-G-sp-G-sp-G-sp-G-sp-G) \ \left(9CI\right)$ CN (CA INDEX NAME) *** STRUCTURE DIAGRAM IS NOT AVAILABLE *** ANSWER 19 OF 21 HCAPLUS COPYRIGHT 2004 ACS on STN L21 2001:125171 HCAPLUS DN 134:294478 Entered STN: 21 Feb 2001 ED TT Human peripheral blood cells differentially recognize and respond to two distinct CpG motifs ΑIJ Verthelyi, Daniela; Ishii, Ken J.; Gursel, Mayda; Takeshita, Fumihiko; Klinman, Dennis M. Section of Retroviral Research, Center for Biologics Evaluation and CS Research, Food and Drug Administration, Bethesda, MD, 20892, USA Journal of Immunology (2001), 166(4), 2372-2377 CODEN: JOIMA3; ISSN: 0022-1767 PB American Association of Immunologists DT Journal English CC 15-10 (Immunochemistry) Oligodeoxynucleotides (ODN) that contain unmethylated CpG dinucleotides AB trigger a strong innate immune response in vertebrates. CpG ODN show promise as vaccine adjuvants, anti-allergens, and immunoprotective agents in animal models. Their transition to clin. use requires the identification of motifs that are optimally stimulatory in humans. Anal. of hundreds of novel ODN resulted in the identification and characterization of two structurally distinct "clusters" of immunostimulatory CpG ODN. One cluster ("D") preferentially stimulates IFN- gamma. production by NK cells, whereas the other ("K") stimulates cell proliferation and the production of IL-6 and IgM by monocytes and B cells. The distinct immunostimulatory properties of K and D ODN can improve the design of CpG-based products to achieve specific therapeutic goals. CpG motif vaccine monocyte lymphocyte cytokine IgM Genetic element RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); PRP (Properties); BIOL (Biological study) (CpG island; two distinct CpG motifs in stimulation of human peripheral blood cells) IT Immunoglobulins RL: BOC (Biological occurrence); BSU (Biological study, unclassified); BIOL (Biological study); OCCU (Occurrence) (M; two distinct CpG motifs in stimulation of human peripheral blood cells) TT Immunostimulants (adjuvants; two distinct CpG motifs in stimulation of human peripheral blood cells) IT Lymphocyte (natural killer cell; two distinct CpG motifs in stimulation of human peripheral blood cells) Allergy IT B cell (lymphocyte) Cell proliferation

```
Monocyte
     Vaccines
        (two distinct CpG motifs in stimulation of human peripheral blood
        cells)
IT
     Oligodeoxyribonucleotides
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); PRP (Properties); BIOL (Biological study)
        (two distinct CpG motifs in stimulation of human peripheral blood
        cells)
     Interleukin 6
IT
     RL: BOC (Biological occurrence); BSU (Biological study, unclassified);
     BIOL (Biological study); OCCU (Occurrence)
        (two distinct CpG motifs in stimulation of human peripheral blood
        cells)
TT
     Interferons
     RL: BOC (Biological occurrence); BSU (Biological study, unclassified);
     BIOL (Biological study); OCCU (Occurrence)
        (.gamma.; two distinct CpG motifs in stimulation of human peripheral
        blood cells)
IT
     246163-61-1 334074-91-8 334074-92-9
                                             334581-75-8
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); BIOL (Biological study)
        (two distinct CpG motifs in stimulation of human peripheral blood
        cells)
RE.CNT 28
              THERE ARE 28 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE
(1) Ballas, Z; J Immunol 1996, V157, P1840 HCAPLUS(2) Bauer, M; Immunology 1999, V97, P699 HCAPLUS
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(19) Pisetsky, D; Immunol Res 1999, V19, P35 HCAPLUS
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     334074-91-8
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); BIOL (Biological study)
        (two distinct CpG motifs in stimulation of human peripheral blood
        cells)
RN
     334074-91-8 HCAPLUS
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     (CA INDEX NAME)
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
L21
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AN
     2001:31630 HCAPLUS
DN
     134:96243
     Entered STN: 12 Jan 2001
ED
     DNA-binding protein CTCF and insulators sequences having specific
ΤI
     enhancer-blocking activity for regulation of gene expression
TN
     Bell, Adam C.; West, Adam G.; Felsenfeld, Gary
     United States Dept. of Health and Human
PA
     Services, USA
so
     PCT Int. Appl., 96 pp.
     CODEN: PIXXD2
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DΤ
     Patent
     English
LΑ
     ICM C12N015-00
IC
     3-2 (Biochemical Genetics)
     Section cross-reference(s): 12, 13
FAN. CNT 1
                                             APPLICATION NO.
                                                                     DATE
     PATENT NO.
                         KIND DATE
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             IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV,
             MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG,
             SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
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PRAI US 1999-141728P
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                                 19990630
    WO 2000-US10509
                                 20000419
CLASS
                 CLASS PATENT FAMILY CLASSIFICATION CODES
PATENT NO.
WO 2001002553 ICM C12N015-00
     An insulator element shown to be both necessary and sufficient for the
     enhancer-blocking activity in human cells is described. The insulator
     element of the invention is a 42-bp DNA mol. having specific enhancer
     blocking activity for regulation of gene expression and found in the chicken .beta.-globin gene cluster. The insulator element has also been
     shown to be the core binding site for CTCF, a DNA binding protein that is
     highly conserved in vertebrates. In addition, an insulator element containing
     CTCF binding sites has been identified in the region of the mouse, rat,
     and human insulin-like growth factor 2 Igf2 locus which is methylated
     exclusively on the paternal allele. Methylation of the insulator sequence
     abolishes the ability of CTCF to bind to the insulator and results in loss
     of enhancer-blocking activity.
    enhancer blocking insulator element gene expression; CTCF DNA binding
     protein insulator element; globin gene insulator element enhancer
     blocking; Igf2 gene insulator element enhancer blocking
    Antibiotic resistance
TТ
     Drugs
     Molecular cloning
        (DNA-binding protein CTCF and insulators sequences having specific
        enhancer-blocking activity for regulation of gene expression)
IT
     Enzymes, preparation
     Hormones, animal, preparation
     Proteins, general, preparation
     RL: BPN (Biosynthetic preparation); BIOL (Biological study); PREP
     (Preparation)
        (DNA-binding protein CTCF and insulators sequences having specific
        enhancer-blocking activity for regulation of gene expression)
IT
     Enhancer (genetic element)
     RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL
     (Biological study); PROC (Process)
        (DNA-binding protein CTCF and insulators sequences having specific
        enhancer-blocking activity for regulation of gene expression)
IT
     Proteins, specific or class
     RL: BPR (Biological process); BSU (Biological study, unclassified); BUU
     (Biological use, unclassified); BIOL (Biological study); PROC (Process);
        (DNA-binding, CTCF; DNA-binding protein CTCF and insulators sequences
        having specific enhancer-blocking activity for regulation of gene
        expression)
IT
     Transcription factors
     RL: BPR (Biological process); BSU (Biological study, unclassified); BUU
     (Biological use, unclassified); BIOL (Biological study); PROC (Process);
     USES (Uses)
        (GAL4; DNA-binding protein CTCF and insulators sequences having
```

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specific enhancer-blocking activity for regulation of gene expression)
TT
     Genetic element
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); BUU (Biological use, unclassified); BIOL (Biological
     study); USES (Uses)
        (ICR (imprinted control region); DNA-binding protein CTCF and
        insulators sequences having specific enhancer-blocking activity for
        regulation of gene expression)
IT
     Methylation
        (biol., insulator element inhibition by; DNA-binding protein CTCF and
        insulators sequences having specific enhancer-blocking activity for
        regulation of gene expression)
TΥ
     Gene
        (expression; DNA-binding protein CTCF and insulators sequences having
        specific enhancer-blocking activity for regulation of gene expression)
     Transcription factors
IT
     RL: BPR (Biological process); BSU (Biological study, unclassified); BUU
     (Biological use, unclassified); BIOL (Biological study); PROC (Process);
     USES (Uses)
        (gene lexA; DNA-binding protein CTCF and insulators sequences having
        specific enhancer-blocking activity for regulation of gene expression)
IT
     Mouse
     Rat
        (insulator element from Igf2 locus of; DNA-binding protein CTCF and
        insulators sequences having specific enhancer-blocking activity for
        regulation of gene expression)
IT
     Hemoglobins
     RL: BSU (Biological study, unclassified); BIOL (Biological study) (insulator element from .beta.-globin locus for; DNA-binding protein
        CTCF and insulators sequences having specific enhancer-blocking
        activity for regulation of gene expression)
     Chicken (Gallus domesticus)
TΤ
        (insulator element from .beta.-globin locus of; DNA-binding protein
        CTCF and insulators sequences having specific enhancer-blocking
        activity for regulation of gene expression)
IT
     Genetic element
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); BUU (Biological use, unclassified); BIOL (Biological
     study); USES (Uses)
        (insulator element; DNA-binding protein CTCF and insulators sequences
        having specific enhancer-blocking activity for regulation of gene
        expression)
     9037-42-7, DNA methyltransferase
     RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
     (Uses)
        (3; DNA-binding protein CTCF and insulators sequences having specific
        enhancer-blocking activity for regulation of gene expression)
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); BUU (Biological use, unclassified); BIOL (Biological
     study); USES (Uses)
        (insulator element consensus from Igf2-H19 locus; DNA-binding protein
        CTCF and insulators sequences having specific enhancer-blocking
        activity for regulation of gene expression)
TT
     319022-93-0 319023-09-1
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); BUU (Biological use, unclassified); BIOL (Biological
     study): USES (Uses)
        (insulator element from chicken .beta.-globin locus; DNA-binding
        protein CTCF and insulators sequences having specific enhancer-blocking
        activity for regulation of gene expression)
     67763-97-7, Insulin-like growth factor 2
IT
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (insulator element from gene for; DNA-binding protein CTCF and
        insulators sequences having specific enhancer-blocking activity for
        regulation of gene expression)
                  319023-03-5
                                319023-04-6
IT
     319023-02-4
                                               319023-05-7 319023-06-8
     319023-07-9
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); BUU (Biological use, unclassified); BIOL (Biological
     study); USES (Uses)
        (insulator element from human Igf2-H19 locus; DNA-binding protein CTCF
        and insulators sequences having specific enhancer-blocking activity for
        regulation of gene expression)
IT
     319022-94-1 319022-95-2 319022-96-3
                                                319022-97-4
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
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study, unclassified); BUU (Biological use, unclassified); BIOL (Biological
     study); USES (Uses)
        (insulator element from mouse Igf2-H19 locus; DNA-binding protein CTCF
        and insulators sequences having specific enhancer-blocking activity for
        regulation of gene expression)
IT
     319022-98-5 319022-99-6 319023-00-2
                                              319023-01-3
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     study, unclassified); BUU (Biological use, unclassified); BIOL (Biological
     study); USES (Uses)
        (insulator element from rat Igf2-H19 locus; DNA-binding protein CTCF
        and insulators sequences having specific enhancer-blocking activity for
        regulation of gene expression)
IT
     1404-04-2, Neomycin 6379-56-2, Hygromycin
     RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL
     (Biological study); PROC (Process)
        (resistance to; DNA-binding protein CTCF and insulators sequences
        having specific enhancer-blocking activity for regulation of gene
        expression)
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     318409-01-7
                  318409-02-8
                                               318409-04-0
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     RL: PRP (Properties)
        (unclaimed nucleotide sequence; dNA-binding protein CTCF and insulators
        sequences having specific enhancer-blocking activity for regulation of
        gene expression)
IT
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        (unclaimed sequence; dNA-binding protein CTCF and insulators sequences
        having specific enhancer-blocking activity for regulation of gene
        expression)
IT
     319025-35-9
     RL: PRP (Properties)
        (unclaimed nucleotide sequence; dNA-binding protein CTCF and insulators
        sequences having specific enhancer-blocking activity for regulation of
        gene expression)
RN
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     35: PN: WO0102553 SEQID: 36 unclaimed DNA (9CI) (CA INDEX NAME)
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*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
L21 ANSWER 21 OF 21 HCAPLUS COPYRIGHT 2004 ACS on STN
     2000:741930 HCAPLUS
AΝ
DN
     133:320986
     Entered STN: 20 Oct 2000
     Oligodeoxynucleotide and its use to induce an immune response
ΤI
IN
     Klinman, Dennis; Ishii, Ken; Verthelyi,
     Daniela
PΑ
     United States Dept. of Health and Human
     Services, USA
PCT Int. Appl., 46 pp.
so
     CODEN: PIXXD2
DT
     Patent
     English
LΑ
IC
     ICM A61K031-70
     ICS A61K039-39; C07H021-00
     15-2 (Immunochemistry)
CC
     Section cross-reference(s): 3, 63
FAN.CNT 2
                         KIND
                              DATE
                                            APPLICATION NO.
     PATENT NO.
                                                                   DATE
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20001019
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     WO 2000061151
                           A3
                                  20010426
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             LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA,
             ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
         RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF,
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                                              EP 2000-923283
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PRAI US 1999-128898P
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                                  19990412
     WO 2000-US9839
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CLASS
                  CLASS PATENT FAMILY CLASSIFICATION CODES
PATENT NO.
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                 ICM
                         A61K031-70
 WO 2000061151
                  ICS
                         A61K039-39; C07H021-00
     The present invention provides a substantially pure or isolated
AB
     oligodeoxynucleotide of at least about 10 nucleotides comprising a
     sequence represented by either the formula: 5' N1N2N3T-CpG-WN4N5N6 3'
     wherein the central CpG motif is unmethylated, W is A or T, and N1, N2,
     N3, N4, N5, and N6 are any nucleotides, or the formula: 5' RY-CpG-RY 3'
     wherein the central CpG motif is unmethylated, R is A or G, and Y is C or
     T, as well as an oligodeoxynucleotide delivery complex and a pharmacol.
     composition comprising the present inventive oligodeoxynucleotide, and a method
     of inducing an immune response by administering the present inventive oligodeoxynucleotide to a host. The oligodeoxynucleotides with phosphate
     or phosphorothicate backbone modification are useful for inducing
     cell-mediated and humoral immune response and are therefore useful for
     treatment of allergy, asthma, cancer, autoimmune disease, immunol.
     disease, infection, and immune deficiency.
     oligodeoxynucleotide adjuvant vaccine allergy cancer infection; autoimmune
ST
     disease asthma oligodeoxynucleotide adjuvant vaccine
IT
     Cell activation
        (B cell; oligodeoxynucleotide for use to induce immune response)
TT
     B cell (lymphocyte)
     Lymphocyte
        (activation; oligodeoxynucleotide for use to induce immune response)
тт
     Infection
        (anti-; oligodeoxynucleotide for use to induce immune response)
TT
     Chemical warfare agents
         (bio-; oligodeoxynucleotide for use to induce immune response)
IT
     Diagnosis
        (cancer; oligodeoxynucleotide for use to induce immune response)
IT
     Drug delivery systems
         (carriers; oligodeoxynucleotide for use to induce immune response)
     Immunity
IT
        (cell-mediated; oligodeoxynucleotide for use to induce immune response)
IT
     Neoplasm
        (diagnosis; oligodeoxynucleotide for use to induce immune response)
IT
        (disorder; oligodeoxynucleotide for use to induce immune response)
IT
     Immunity
        (humoral; oligodeoxynucleotide for use to induce immune response)
IT
     Drug delivery systems
        (liposomes; oligodeoxynucleotide for use to induce immune response)
IT
     Cell activation
        (lymphocyte; oligodeoxynucleotide for use to induce immune response)
IT
     Animal cell
        (non-B; oligodeoxynucleotide for use to induce immune response)
     Allergy
     Allergy inhibitors
     Antitumor agents
     Asthma
     Autoimmune disease
     B cell (lymphocyte)
     Drug targeting
     Immunodeficiency
     Immunostimulants
     Neoplasm
     Vaccines
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(oligodeoxynucleotide for use to induce immune response)
IT
    Antibodies
     Cytokines
     Interleukin 6
     RL: BOC (Biological occurrence); BSU (Biological study, unclassified); MFM
     (Metabolic formation); THU (Therapeutic use); BIOL (Biological study);
     FORM (Formation, nonpreparative); OCCU (Occurrence); USES (Uses)
        (oligodeoxynucleotide for use to induce immune response)
     Lipids, biological studies
     Oligodeoxyribonucleotides
    RL: BSU (Biological study, unclassified); PRP (Properties); THU
     (Therapeutic use); BIOL (Biological study); USES (Uses)
        (oligodeoxynucleotide for use to induce immune response)
IT
    Neoplasm
        (solid; oligodeoxynucleotide for use to induce immune response)
IT
    Antisense oligonucleotides
     RL: BOC (Biological occurrence); BSU (Biological study, unclassified); MFM
     (Metabolic formation); THU (Therapeutic use); BIOL (Biological study);
     FORM (Formation, nonpreparative); OCCU (Occurrence); USES (Uses)
        (therapy; oligodeoxynucleotide for use to induce immune response)
ΙT
        (virosomes; oligodeoxynucleotide for use to induce immune response)
IT
     Interferons
    RL: BOC (Biological occurrence); BSU (Biological study, unclassified); MFM
     (Metabolic formation); THU (Therapeutic use); BIOL (Biological study);
     FORM (Formation, nonpreparative); OCCU (Occurrence); USES (Uses)
        (.gamma.; oligodeoxynucleotide for use to induce immune response)
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301939-87-7D,

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57-88-5, Cholesterol, biological studies
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AN
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    Oligonucleotide probes in nucleic acid arrays for genetic analysis of
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IN
    Mittman, Michael; Mack, David J.; Lockhart, David J.
PA
    Affymetrix, Inc., USA
    U.S., 183 pp.
CODEN: USXXAM
SO
DT
    Patent
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    English
    C12Q001-68; C07H021-02; C07H021-04; G01N015-06; G01N033-00
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    435006000; 422068100; 536023100; 536024300; 536024310
NCL
     3-1 (Biochemical Genetics)
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                       435006000; 422068100; 536023100; 536024300; 536024310
    This invention provides 127,806 unique oligonucleotide sequences which are
AB
     complementary to .apprx.6500 specific known murine genes. The invention
    provides the sequences in such a way as to make them available for gene
     expression monitoring by hybridization to high d. oligonucleotide arrays,
     or for use as primers for PCR and other amplification protocols. As such,
     the invention related to diverse fields impacted by the nature of mol.
     interaction, including chemical, biol., medicine, and medical diagnostics.
     [This abstract record is one of 26 records for this document necessitated by
     the large number of index entries required to fully index the document and
     publication system constraints.].
ST
    oligonucleotide probe microarray genetic analysis mouse; gene expression
     mouse oligonucleotide probe hybridization
    DNA microarray technology
     DNA sequences
    Gene expression profiles, animal
     Genetic methods
     Mus musculus
    Nucleic acid amplification (method)
    Nucleic acid hybridization
     PCR (polymerase chain reaction)
        (oligonucleotide probes in nucleic acid arrays for genetic anal. of
       mouse)
TT
    Primers (nucleic acid)
     Probes (nucleic acid)
     RL: ARG (Analytical reagent use); PRP (Properties); ANST (Analytical
     study); USES (Uses)
        (oligonucleotide probes in nucleic acid arrays for genetic anal. of
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    RL: ARG (Analytical reagent use); PRP (Properties); ANST (Analytical
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795709-39-6

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(probe and/or primer; oligonucleotide probes in nucleic acid arrays for
        genetic anal. of mouse)
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        (probe and/or primer; oligonucleotide probes in nucleic acid arrays for
        genetic anal. of mouse)
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RL: ARG (Analytical reagent use); PRP (Properties); ANST (Analytical
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     RL: ARG (Analytical reagent use); PRP (Properties); ANST (Analytical
     study); USES (Uses)
        (probe and/or primer; oligonucleotide probes in nucleic acid arrays for
        genetic anal. of mouse)
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L27
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    2004:1022684 HCAPLUS
Entered STN: 29 Nov 2004
ΔN
ED
    Oligonucleotide probes in nucleic acid arrays for genetic analysis of
ΤI
    mouse
    Mittman, Michael; Mack, David J.; Lockhart, David J.
IN
PA
    Affymetrix, Inc., USA
so
    U.S., 183 pp.
    CODEN: USXXAM
DT
    Patent
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    English
IC
    C12Q001-68; C07H021-02; C07H021-04; G01N015-06; G01N033-00
    435006000; 422068100; 536023100; 536024300; 536024310
NCL
    3-1 (Biochemical Genetics)
CC
    Section cross-reference(s): 13
FAN.CNT 26
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CLASS
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                NCL
    This invention provides 127,806 unique oligonucleotide sequences which are
AB
    complementary to .apprx.6500 specific known murine genes. The invention
    provides the sequences in such a way as to make them available for gene
    expression monitoring by hybridization to high d. oligonucleotide arrays,
    or for use as primers for PCR and other amplification protocols. As such,
    the invention related to diverse fields impacted by the nature of mol.
     interaction, including chemical, biol., medicine, and medical diagnostics.
     [This abstract record is one of 26 records for this document necessitated by
    the large number of index entries required to fully index the document and
    publication system constraints.].
    oligonucleotide probe microarray genetic analysis mouse; gene expression
ST
    mouse oligonucleotide probe hybridization
IT
    DNA microarray technology
    DNA sequences
    Gene expression profiles, animal
    Genetic methods
    Mus musculus
    Nucleic acid amplification (method)
    Nucleic acid hybridization
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PCR (polymerase chain reaction)

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(oligonucleotide probes in nucleic acid arrays for genetic anal. of
        mouse)
     Primers (nucleic acid)
IT
     Probes (nucleic acid)
     RL: ARG (Analytical reagent use); PRP (Properties); ANST (Analytical
     study); USES (Uses)
        (oligonucleotide probes in nucleic acid arrays for genetic anal. of
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genetic anal. of mouse)

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795596-30-4

795596-32-6

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RL: ARG (Analytical reagent use); PRP (Properties); ANST (Analytical
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Entered STN: 29 Nov 2004
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    Mittman, Michael; Mack, David J.; Lockhart, David J.
IN
    Affymetrix, Inc., USA
PA
    U.S., 183 pp.
CODEN: USXXAM
SO
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LΑ
    English
    C12Q001-68; C07H021-02; C07H021-04; G01N015-06; G01N033-00
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    complementary to .apprx.6500 specific known murine genes. The invention
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AB This invention provides 127,806 unique oligonucleotide sequences which are complementary to .apprx.6500 specific known murine genes. The invention provides the sequences in such a way as to make them available for gene expression monitoring by hybridization to high d. oligonucleotide arrays, or for use as primers for PCR and other amplification protocols. As such, the invention related to diverse fields impacted by the nature of mol. interaction, including chemical, biol., medicine, and medical diagnostics. [This abstract record is one of 26 records for this document necessitated by the large number of index entries required to fully index the document and

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publication system constraints.].
ST
     oligonucleotide probe microarray genetic analysis mouse; gene expression
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ΙT
    DNA microarray technology
    DNA sequences
    Gene expression profiles, animal
     Genetic methods
     Mus musculus
    Nucleic acid amplification (method)
    Nucleic acid hybridization
     PCR (polymerase chain reaction)
        (oligonucleotide probes in nucleic acid arrays for genetic anal. of
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        (probe and/or primer; oligonucleotide probes in nucleic acid arrays for
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   (probe and/or primer; oligonucleotide probes in nucleic acid arrays for
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     Mittman, Michael; Mack, David J.; Lockhart, David J.
     Affymetrix, Inc., USA
     U.S., 183 pp.
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     This invention provides 127,806 unique oligonucleotide sequences which are
     complementary to .apprx.6500 specific known murine genes. The invention
     provides the sequences in such a way as to make them available for gene
     expression monitoring by hybridization to high d. oligonucleotide arrays,
     or for use as primers for PCR and other amplification protocols. As such,
     the invention related to diverse fields impacted by the nature of mol.
     interaction, including chemical, biol., medicine, and medical diagnostics. [This abstract record is one of 26 records for this document necessitated by
     the large number of index entries required to fully index the document and
     publication system constraints.].
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     DNA microarray technology
     DNA sequences
     Gene expression profiles, animal
     Genetic methods
     Mus musculus
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        (oligonucleotide probes in nucleic acid arrays for genetic anal. of
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study); USES (Uses)
   (probe and/or primer; oligonucleotide probes in nucleic acid arrays for
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     The present invention concerns methods and reagents useful in modulating
     hepatitis C virus (HCV) gene expression in a variety of applications,
     including use in therapeutic, diagnostic, target validation, and genomic
     discovery applications. Specifically, the invention relates to small
     nucleic acid mols., such as short interfering nucleic acid (siNA), short
     interfering RNA (siRNA), double-stranded RNA (dsRNA), micro-RNA (miRNA),
     and short hairpin RNA (shRNA) mols. capable of mediating RNA interference
     (RNAi) against hepatitis C virus (HCV) gene expression and/or activity.
     The small nucleic acid mols. are useful in the treatment and diagnosis of
     HCV infection, liver failure, hepatocellular carcinoma, cirrhosis and any
     other disease or condition that responds to modulation of HCV expression
     or activity.
ST
     RNA interference inhibition gene hepatitis virus HCV
IT
     Pyrimidine nucleotides
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (2'-O-Me, 2'-deoxy-2'-fluoro; RNA interference mediated inhibition of
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IT
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        (2'-deoxy, 2'-0-Me; RNA interference mediated inhibition of hepatitis C
        virus (HCV) gene expression using short interfering nucleic acid
        (sina))
IT
     Gene, microbial
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RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (HCV, expression; RNA interference mediated inhibition of hepatitis C
        virus (HCV) gene expression using short interfering nucleic acid
        (sina))
IT
     Antiviral agents
     Hepatitis C virus
     Human
        (RNA interference mediated inhibition of hepatitis C virus (HCV) gene
        expression using short interfering nucleic acid (sina))
     Antisense nucleic acids
TT
     Oligonucleotides
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (RNA interference mediated inhibition of hepatitis C virus (HCV) gene
        expression using short interfering nucleic acid (sina))
TΤ
     Post-transcriptional processing
        (interference; RNA interference mediated inhibition of hepatitis C
        virus (HCV) gene expression using short interfering nucleic acid
        (sina))
IT
     Nucleic acids
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (short interfering; RNA interference mediated inhibition of hepatitis C
        virus (HCV) gene expression using short interfering nucleic acid
        (sina))
TT
     Double stranded RNA
     RL: BPN (Biosynthetic preparation); THU (Therapeutic use); BIOL
     (Biological study); PREP (Preparation); USES (Uses)
        (small interfering; RNA interference mediated inhibition of hepatitis C
        virus (HCV) gene expression using short interfering nucleic acid
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        C virus (HCV) gene expression using short interfering nucleic acid
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        (unclaimed sequence; rNA interference mediated inhibition of hepatitis
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        C virus (HCV) gene expression using short interfering nucleic acid
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        (unclaimed sequence; rNA interference mediated inhibition of hepatitis
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     RL: PRP (Properties)
        (unclaimed sequence; rNA interference mediated inhibition of hepatitis
        C virus (HCV) gene expression using short interfering nucleic acid
        (sina))
IT
     777957-59-2 777958-74-4
     RL: PRP (Properties)
        (unclaimed sequence; rNA interference mediated inhibition of hepatitis
        C virus (HCV) gene expression using short interfering nucleic acid
        (sina))
RN
     777957-59-2 HCAPLUS
     RNA, (U-U-C-G-C-C-G-A-C-C-U-C-A-U-G-G-G-U) (9CI) (CA INDEX NAME)
CN
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
     777958-74-4 HCAPLUS
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     RNA, (C-U-U-C-G-C-C-G-A-C-C-U-C-A-U-G-G-G-G) (9CI) (CA INDEX NAME)
CN
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
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L27
     2004:817415 HCAPLUS
AN
DN
     141:309637
ED
     Entered STN: 07 Oct 2004
     Transgenic plants expressing chymotrypsin inhibitor mutant proteins with
ΤI
     increased levels of essential amino acids for food or feed use
TN
    Rao, Aragula Gururaj; Roesler, Keith R.
    Pioneer Hi-Bred International, Inc., USA
U.S., 47 pp., Cont.-in-part of U.S. Ser. No. 297,418, abandoned.
PA
     CODEN: USXXAM
DT
    Patent
LΑ
    English
IC
     ICM C07K014-00
    530300000; 530350000; 530370000; 426044000
     7-3 (Enzymes)
CC
     Section cross-reference(s): 3, 11, 17
FAN.CNT 2
    PATENT NO.
                         KIND
                                DATE
                                             APPLICATION NO.
                                                                     DATE
                                20041005 US 1999-311689
    US 6800726
                         В1
                                                                     19990513 <--
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PRAI US 1996-740682
                          B2
                                19961101 <--
    US 1999-297418
                          B2
                                19990420 <--
CLASS
                 CLASS PATENT FAMILY CLASSIFICATION CODES
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                        C07K014-00
                 NCL
                        530300000; 530350000; 530370000; 426044000
                        A23K001/16G; A23L001/305C; C07K014/81; C07K014/81B1;
US 6800726
                 ECLA
                        C12N015/82C4; C12N015/82C4B10; C12N015/82C4B10A;
                        C12N015/82C4B10B
    The present invention provides methods for increasing essential amino acid
     content of plants using modified polypeptides derived from a protease
     inhibitor, and more specifically, a chymotrypsin inhibitor. The invention
     based on the discovery that chymotrypsin inhibitor-2 (CI-2) can be
     modified to enhance its essential amino acid content. Protease inhibitor
     proteins can be modified from the proteins of barley or maize by
     substitution mutation to increase the essential amino acid content and
     decrease protease inhibitor activities. Sequences for wild-type CI-2
     protein from barley, as well as truncated and mutated variants, are
     provided. Other plant species that may be used as a source for
     chymotrypsin inhibitor include Zea mays, Vicia faba, Cucurbita maxima,
     Canavalia lineata, Vigna angularis, Nicotiana tabacum, Nicotiana
sylvestris, Sambucus nigra, Momordica charantia, Solanum tuberosum,
     Lycopersicon peruvianum, Lycopersicon esculentum, Amaranthus caudatus and
     Arabidopsis thaliana. The invention further provides recombinant expression cassettes, host cells, transgenic plants, and antibody compns.
     The compds. of the invention are thus excellent candidates for
     transformation of feed grain and food to enhance nutrition.
    plant protease inhibitor mutant feed essential amino acid; chymotrypsin
     inhibitor essential amino acid feed transgenic plant; barley chymotrypsin
     inhibitor essential amino acid mutant; proteinase inhibitor CI2 sequence
     homolog cDNA Hordeum
     Fusion proteins (chimeric proteins)
IT
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RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     BIOL (Biological study); PREP (Preparation)
        (BHL3N dimer; transgenic plants expressing chymotrypsin inhibitor
        mutant proteins with increased levels of essential amino acids for food
        or feed use)
IT
     Sequence homology analysis
        (BLAST, use of; transgenic plants expressing chymotrypsin inhibitor
        mutant proteins with increased levels of essential amino acids for food
        or feed use)
    Amaranthus caudatus
IT
     Arabidopsis thaliana
     Canavalia lineata
     Cucurbita maxima
     Hordeum vulgare
     Lycopersicon esculentum
     Lycopersicon peruvianum
     Momordica charantia
     Nicotiana sylvestris
     Nicotiana tabacum
     Sambucus nigra
     Solanum tuberosum
     Vicia faba
     Vigna angularis
     Zea mays
        (chymotrypsin inhibitor from; transgenic plants expressing chymotrypsin
        inhibitor mutant proteins with increased levels of essential amino
        acids for food or feed use)
IT
     Amino acids, biological studies
     RL: BPN (Biosynthetic preparation); FFD (Food or feed use); BIOL
     (Biological study); PREP (Preparation); USES (Uses)
        (essential; transgenic plants expressing chymotrypsin inhibitor mutant
        proteins with increased levels of essential amino acids for food or
IT
     Digestion, biological
        (improved digestibiluty of engineered proteins; transgenic plants
        expressing chymotrypsin inhibitor mutant proteins with increased levels
        of essential amino acids for food or feed use)
IT
     Disulfide group
        (increased stability achieved by addition of; transgenic plants expressing
        chymotrypsin inhibitor mutant proteins with increased levels of
        essential amino acids for food or feed use)
     Mutagenesis
        (site-directed, substitution; transgenic plants expressing chymotrypsin
        inhibitor mutant proteins with increased levels of essential amino
        acids for food or feed use)
     Antibodies and Immunoglobulins
     RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
     (Uses)
        (to epitopes of protease inhibitors; transgenic plants expressing
        chymotrypsin inhibitor mutant proteins with increased levels of
        essential amino acids for food or feed use)
TT
     Feed
     Food
     Genetic engineering
     Molecular cloning
     Protein engineering
     Protein sequences
     cDNA sequences
        (transgenic plants expressing chymotrypsin inhibitor mutant proteins
        with increased levels of essential amino acids for food or feed use)
IT
     Embryophyta
        (transgenic; transgenic plants expressing chymotrypsin inhibitor mutant
        proteins with increased levels of essential amino acids for food or
IT
     56-87-1P, Lysine, biological studies
     RL: BPN (Biosynthetic preparation); FFD (Food or feed use); BIOL
     (Biological study); PREP (Preparation); USES (Uses)
        (-high content variants; transgenic plants expressing chymotrypsin
        inhibitor mutant proteins with increased levels of essential amino
        acids for food or feed use)
    765830-30-6DP, mutated variants
                                        765830-32-8P
IT
                                                       765830-34-0P
     765830-36-2P
                    765830-38-4P
                                   765830-40-8P
                                                  765830-42-0P
                    765830-48-6P
     765830-46-4P
     RL: AGR (Agricultural use); BPN (Biosynthetic preparation); BSU
     (Biological study, unclassified); FFD (Food or feed use); PRP (Properties); BIOL (Biological study); PREP (Preparation); USES (Uses)
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(amino acid sequence; transgenic plants expressing chymotrypsin
        inhibitor mutant proteins with increased levels of essential amino
        acids for food or feed use)
    12063-98-8, Gallium phosphide, analysis
TT
     RL: ARU (Analytical role, unclassified); ANST (Analytical study)
        (anal.; transgenic plants expressing chymotrypsin inhibitor mutant
        proteins with increased levels of essential amino acids for food or
        feed use)
     765830-29-3D, mutated variants
                                      765830-31-7
                                                    765830-33-9
IT
                                                            765830-45-3
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     765830-37-3
                  765830-39-5
     765830-47-5
     RL: AGR (Agricultural use); BSU (Biological study, unclassified); BUU
     (Biological use, unclassified); PRP (Properties); BIOL (Biological study);
     USES (Uses)
        (nucleotide sequence; transgenic plants expressing chymotrypsin
        inhibitor mutant proteins with increased levels of essential amino
        acids for food or feed use)
TT
    139466-47-0DP, Chymotrypsin inhibitor 2, derivs.
     RL: AGR (Agricultural use); BPN (Biosynthetic preparation); BSU
     (Biological study, unclassified); FFD (Food or feed use); PRP
     (Properties); BIOL (Biological study); PREP (Preparation); USES (Uses)
        (transgenic plants expressing chymotrypsin inhibitor mutant proteins
        with increased levels of essential amino acids for food or feed use)
     37205-61-1D, Protease inhibitor, derivs.
                                               141256-43-1D, Chymotrypsin
     inhibitor, derivs.
    RL: AGR (Agricultural use); BSU (Biological study, unclassified); FFD (Food or feed use); BIOL (Biological study); USES (Uses)
        (transgenic plants expressing chymotrypsin inhibitor mutant proteins
        with increased levels of essential amino acids for food or feed use)
     63-68-3P, Methionine, biological studies 72-19-5P, Threonine, biological
IT
               73-22-3P, Tryptophan, biological studies
     studies
     RL: BPN (Biosynthetic preparation); FFD (Food or feed use); BIOL
     (Biological study); PREP (Preparation); USES (Uses)
        (transgenic plants expressing chymotrypsin inhibitor mutant proteins
        with increased levels of essential amino acids for food or feed use)
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     765830-99-7
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                                765831-01-4
                                               765831-02-5
     RL: PRP (Properties)
        (unclaimed nucleotide sequence; transgenic plants expressing
        chymotrypsin inhibitor mutant proteins with increased levels of
        essential amino acids for food or feed use)
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                                 765830-77-1
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     765830-80-6
     RL: PRP (Properties)
        (unclaimed protein sequence; transgenic plants expressing chymotrypsin
        inhibitor mutant proteins with increased levels of essential amino
        acids for food or feed use)
              THERE ARE 26 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT
RE
(1) Anon; WO 9007862 1990
(2) Anon; EP 0485970 WO 1992 HCAPLUS
(3) Anon; WO 9205239 1992 HCAPLUS
(4) Anon; WO 9320175 1993 HCAPLUS
(5) Anon; WO 9410315 1994 HCAPLUS
(6) Anon; WO 9416078 1994 HCAPLUS
(7) Anon; WO 9527068 1995 HCAPLUS
(8) Burow; Mol Gen Genet 1993, V241, P431 HCAPLUS
(9) Campbell, A; Plant Protein Engineering 1992, P257 HCAPLUS
(10) Clore; Protein Eng 1987, V1, P313 HCAPLUS
(11) Glassman; US 5258300 A 1993 HCAPLUS
(12) Hey; US 6169232 B1 2001 HCAPLUS
(13) Hirabayashi; J Biol Chem 1991, V266, P23648 HCAPLUS
(14) Jaynes; US 5597945 A 1997 HCAPLUS
(15) Jaynes; US 5811654 A 1998 HCAPLUS
(16) Jonassen, I; Carlsberg Res Commun 1980, V45, P47 HCAPLUS
(17) Karchi; The Plant J 1993, V3, P721 HCAPLUS
(18) Lundquist; US 6160208 A 2000
(19) McPhalen; Biochemistry 1987, V26, P261 HCAPLUS
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(20) Mikkelsen; US 5527487 A 1996 HCAPLUS
(21) Mikkelsen; US 5674833 A 1997 HCAPLUS
(22) Ringe, D; Nature 1989, V339, P658 MEDLINE
(23) Summers; J Biol Chem 1989, V264(33), P20082 HCAPLUS
(24) Svendsen, I; Carlsberg Res Commun 1980, V45, P79 HCAPLUS (25) van Eijsden; Plant Mol Biol 1992, V20, P1049 HCAPLUS
(26) Zabin; Biochemistry 1991, V30(25), P6230 HCAPLUS
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     765830-86-2 765830-88-4
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        (unclaimed nucleotide sequence; transgenic plants expressing
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        essential amino acids for food or feed use)
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CN
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CN
     (CA INDEX NAME)
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L27
     2004:780350 HCAPLUS
AN
DN
     141 - 289035
ED
     Entered STN: 24 Sep 2004
     Antisense modulation of CD40 expression for treatment of diseases
IN
     Bennett, C. Frank; Cowsert, Lex M.; Malik, Leila; Siwkowski, Andrew;
     Eldrup, Anne B.
     USA
PΑ
     U.S. Pat. Appl. Publ., 85 pp., Cont.-in-part of WO 2004 29,075.
SO
     CODEN: USXXCO
DT
     Patent
LA
     English
     ICM A61K048-00
IC
     ICS C07H021-04
NCL 514044000; 536023500
    1-7 (Pharmacology)
CC
FAN.CNT 4
     PATENT NO.
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                                           APPLICATION NO.
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                                            US 2003-698689
    US 2004186071
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PΙ
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     US 2002028923
                         A1
                                20020307
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     US 2004063618
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                                                                    20020930
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PRAI US 1998-81483P
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     US 2002-261382
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CT.ASS
PATENT NO.
                 CLASS PATENT FAMILY CLASSIFICATION CODES
                ICM
                        A61K048-00
US 2004186071
                 ICS
                        C07H021-04
                 NCL
                        514044000; 536023500
                 ECLA C12N015/10C4; C12Q001/68A8; G06F019/00A
US 2002028923
   Antisense compds., compns. and methods are provided for inhibiting the
     expression of CD40. The compns. comprise antisense compds., particularly
     antisense oligonucleotides, targeted to nucleic acids encoding CD40.
     Methods of using these compds. for modulation of CD40 expression and for
     treatment of diseases associated with expression of CD40 are provided.
     Antisense oligonucleotides were designed targeting different regions of
     the CD40 mRNA sequence and may be modified to contain phosphorothioate
     linkages, 2'-0-methoxyethyl sugar moiety, and 5-methylcytosine bases. The
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antisense oligonucleotides demonstrated at least 50% inhibition of human CD40 expression. The invention provides methods for synthesis of the antisense oligonucleotides. The antisense oligonucleotides could be used for treatments of hyperproliferative disease, cancer, immune-associated disorder, and inflammatory condition. antisense oligonucleotide human CD40 inhibition; immunomodulator inflammation tumor therapy antisense PNA Human Mus (CD40 from; antisense modulation of CD40 expression for treatment of Interleukin 2 RL: BSU (Biological study, unclassified); BIOL (Biological study) (CD40-dependent production of, antisense reducing; antisense modulation of CD40 expression for treatment of diseases) Disease, animal (CD40-related, treatment of; antisense modulation of CD40 expression for treatment of diseases) Transplant rejection (allotransplant, treatment of; antisense modulation of CD40 expression for treatment of diseases) Antiarteriosclerotics (antiatherosclerotics; antisense modulation of CD40 expression for treatment of diseases) Immunomodulators (antisense as; antisense modulation of CD40 expression for treatment of diseases) Anti-inflammatory agents Antiasthmatics Antirheumatic agents Antitumor agents Drug screening Test kits (antisense modulation of CD40 expression for treatment of diseases) CD40 (antigen) RL: BSU (Biological study, unclassified); BIOL (Biological study) (antisense modulation of CD40 expression for treatment of diseases) Antisense oligonucleotides Phosphorothicate oligonucleotides RL: PAC (Pharmacological activity); PRP (Properties); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses) (antisense modulation of CD40 expression for treatment of diseases) Peptide nucleic acids RL: PAC (Pharmacological activity); PRP (Properties); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses) (antisense; antisense peptide nucleic acid modulation of CD40 expression for treatment of diseases) Immunity (disorder, treatment of; antisense modulation of CD40 expression for treatment of diseases) Transplant and Transplantation (graft-vs.-host reaction, treatment of; antisense modulation of CD40 expression for treatment of diseases) Intestine, disease (inflammatory, treatment of; antisense modulation of CD40 expression for treatment of diseases) B cell (lymphocyte) Macrophage (inhibiting CD 40 in; antisense modulation of CD40 expression for treatment of diseases) RNA splicing (of CD40, antisense redirection of; antisense modulation of CD40 expression for treatment of diseases) Signal transduction, biological (of CD40, antisense reducing; antisense modulation of CD40 expression for treatment of diseases) Diagnosis (of CD40-related disease; antisense modulation of CD40 expression for treatment of diseases)

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Disease, animal

Atherosclerosis

Asthma

expression for treatment of diseases)

Search done by Noble Jarrell

(proliferative, hyper-, treatment of; antisense modulation of CD40

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Autoimmune disease
Inflammation
Neoplasm
Psoriasis
Rheumatoid arthritis
   (treatment of; antisense modulation of CD40 expression for treatment of
   diseases)
Primers (nucleic acid)
Probes (nucleic acid)
RL: ARG (Analytical reagent use); DGN (Diagnostic use); PRP (Properties);
ANST (Analytical study); BIOL (Biological study); USES (Uses)
   (used for diagnosis of CD 40-related disease; antisense modulation of
   CD40 expression for treatment of diseases)
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RL: PAC (Pharmacological activity); PRP (Properties); SPN (Synthetic
preparation); THU (Therapeutic use); BIOL (Biological study); PREP
(Preparation); USES (Uses)
   (antisense modulation of CD40 expression for treatment of diseases)
56-87-1, L-Lysine, biological studies 70-26-8, L-Ornithine
L-Histidine, biological studies 74-79-3, L-Arginine, biological studies
156-86-5, L-Homoarginine 305-62-4, 2,4-Diamino butyric acid 348-66-3, D-Ornithine 351-50-8, D-Histidine 498-56-6, Homolysine 923-27-3, D-Lysine 2259-86-1 4299-56-3, beta.-Lysine 14191-90-3,
L-Norarginine 110798-13-5, D-Homoarginine 760975-35-7 760975-36-8
RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
(Biological study); USES (Uses)
   (in antisense PNA; antisense peptide nucleic acid modulation of CD40
   expression for treatment of diseases)
554-01-8, 5-Methylcytosine
RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
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   (in antisense oligo; antisense modulation of CD40 expression for
   treatment of diseases)
762312-02-7
              762312-03-8
RL: ARG (Analytical reagent use); DGN (Diagnostic use); PRP (Properties);
ANST (Analytical study); BIOL (Biological study); USES (Uses)
   (primer used for diagnosis of CD 40-related disease; antisense
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modulation of CD40 expression for treatment of diseases)
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     RL: ARG (Analytical reagent use); DGN (Diagnostic use); PRP (Properties);
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        (probe used for diagnosis of CD 40-related disease; antisense
        modulation of CD40 expression for treatment of diseases)
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        (unclaimed nucleotide sequence; antisense modulation of CD40 expression
        for treatment of diseases)
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     RL: PRP (Properties)
        (unclaimed sequence; antisense modulation of CD40 expression for
        treatment of diseases)
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     RL: PRP (Properties)
        (unclaimed nucleotide sequence; antisense modulation of CD40 expression
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CN
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
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     140:35886
     Entered STN: 12 Dec 2003
ED
     Identification of genetic targets for modulation by antisense
TI
     oligonucleotides and generation of oligonucleotides for gene modulation
     Cowsert, Lex M.; Baker, Brenda F.; McNeil, John; Freier, Susan M.; Sasmor,
     Henri M.; Brooks, Douglas G.; Ohashi, Cara; Wyatt, Jacqueline R.;
     Borchers, Alexander H.; Vickers, Timothy A.
PA
     USA
SO
     U.S. Pat. Appl. Publ., 344 pp., Cont.-in-part of U.S. Ser. No. 295,463.
     CODEN: USXXCO
DT
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IC
     ICM C12Q001-68
NCL
     435006000
CC
     1-1 (Pharmacology)
     Section cross-reference(s): 3
FAN.CNT 4
                         KIND
                                DATE
                                             APPLICATION NO.
                                                                     DATE
     PATENT NO.
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                ECLA
                       C12N015/10C4; C12Q001/68A8; G06F019/00A
                       C12N015/10C4; C12Q001/68A8; G06F019/00A
US 2002028923 ECLA
    Iterative, preferably computer based iterative processes for generating
     synthetic compds. capable of modulation of target expression are provided.
     During iterations of the processes, a target nucleic acid sequence is
     provided or selected, and a library of candidate nucleobase sequences is
     generated in silico according to defined criteria. A "virtual"
     oligonucleotide chemical is chosen and a library of virtual oligonucleotide
     compds. having the selected nucleobase sequences is generated. These
     virtual compds. are reviewed and compds. predicted to have particular
     properties are selected. The selected compds. are robotically synthesized
     and are preferably robotically assayed for a desired phys., chemical or biol.
     activity. Compds. exhibiting the ability to modulate target expression
     are identified as target modulators. Target modulators thus generated are
     used in assays of parameters indicative of biol. processes to effect gene
     function anal. and in assays of parameters indicative of diseases or
     disorders to effect target valid. The methods are exemplified by
     generation of antisense oligonucleotides targeted to CD40, RhoC, cellular
     inhibitor of apoptosis 2 (c-IAP-2), Gi protein .alpha.-subunit, ELK-1,
     Akt-1, Jagged 2, transforming growth factor .beta.3, apolipoprotein B, BH3
     interacting domain death agonist, and PTEN in the treatment of diseases of
     angiogenesis, breast cancer, and melanoma.
ST
     antisense oligonucleotide generation gene modulation
     Apolipoproteins
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (B, target gene encoding; identification of genetic targets for
        modulation by antisense oligonucleotides and generation of
        oligonucleotides for gene modulation)
TT
     Proteins
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (BH3 interacting domain death agonist, target gene encoding; identification of genetic targets for modulation by antisense
        oligonucleotides and generation of oligonucleotides for gene
        modulation)
     G proteins (guanine nucleotide-binding proteins)
     RL: BSU (Biological study, unclassified); BIOL (Biological study) (Gi (adenylate cyclase-inhibiting), target gene encoding
        .alpha.-subunit of; identification of genetic targets for modulation by
        antisense oligonucleotides and generation of oligonucleotides for gene
        modulation)
IT
     Proteins
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (Jagged 2, target gene encoding; identification of genetic targets for
        modulation by antisense oligonucleotides and generation of
        oligonucleotides for gene modulation)
IT
     Rho protein (G protein)
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (RhoC, target gene encoding; identification of genetic targets for
        modulation by antisense oligonucleotides and generation of
        oligonucleotides for gene modulation)
TΤ
    Angiogenesis
     Apoptosis
     Cell cycle
     Inflammation
        (assay system; identification of genetic targets for modulation by
        antisense oligonucleotides and generation of oligonucleotides for gene
        modulation)
IT
    Glycerides, analysis
     Lipoproteins
     RL: ARU (Analytical role, unclassified); ANST (Analytical study)
        (assay system; identification of genetic targets for modulation by
        antisense oligonucleotides and generation of oligonucleotides for gene
        modulation)
IT
     Proteins
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
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(cIAP-2 (cellular inhibitor of apoptosis protein 2), target gene encoding; identification of genetic targets for modulation by antisense oligonucleotides and generation of oligonucleotides for gene modulation)

IT DNA

RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified); PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(double-stranded; identification of genetic targets for modulation by antisense oligonucleotides and generation of oligonucleotides for gene modulation)

IT Immunoassay

(enzyme-linked immunosorbent assay; identification of genetic targets for modulation by antisense oligonucleotides and generation of oligonucleotides for gene modulation)

IT Hepatitis

(fulminant; identification of genetic targets for modulation by antisense oligonucleotides and generation of oligonucleotides for gene modulation)

IT Disease, animal

(genes related to; identification of genetic targets for modulation by antisense oligonucleotides and generation of oligonucleotides for gene modulation)

IT Angiogenesis inhibitors

Antitumor agents

Computer application

Mammary gland, neoplasm

Melanoma

(identification of genetic targets for modulation by antisense oligonucleotides and generation of oligonucleotides for gene modulation)

IT Antisense oligonucleotides

RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified); PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(identification of genetic targets for modulation by antisense oligonucleotides and generation of oligonucleotides for gene modulation)

IT Gene, animal

RL: BSU (Biological study, unclassified); BIOL (Biological study) (identification of genetic targets for modulation by antisense oligonucleotides and generation of oligonucleotides for gene modulation)

IT Liver, disease

(injury; identification of genetic targets for modulation by antisense oligonucleotides and generation of oligonucleotides for gene modulation)

IT Rodentia

(model test system; identification of genetic targets for modulation by antisense oligonucleotides and generation of oligonucleotides for gene modulation)

IT DNA sequences

(of antisense oligonucleotides targeted for gene modulation)

IT PCR (polymerase chain reaction)

(real-time; identification of genetic targets for modulation by antisense oligonucleotides and generation of oligonucleotides for gene modulation)

IT Double stranded RNA

RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified); PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(small interfering; identification of genetic targets for modulation by antisense oligonucleotides and generation of oligonucleotides for gene modulation)

IT CD40 (antigen)

RL: BSU (Biological study, unclassified); BIOL (Biological study) (target gene encoding; identification of genetic targets for modulation by antisense oligonucleotides and generation of oligonucleotides for gene modulation)

IT Free energy Thermodynamics

(virtual compound selection by evaluation of; identification of genetic targets for modulation by antisense oligonucleotides and generation of oligonucleotides for gene modulation)

IT Transforming growth factors

RL: BSU (Biological study, unclassified); BIOL (Biological study)

(.beta.3-, target gene encoding; identification of genetic targets for modulation by antisense oligonucleotides and generation of oligonucleotides for gene modulation) 404880-85-9, ISIS 119902 404880-86-0, ISIS 120000 404880-87-1, ISIS IT 404880-89-3, ISIS 119845 120001 404880-88-2, ISIS 120002 404880-90-6, ISIS 119846 404880-91-7, ISIS 119847 404880-92-8, ISIS 404880-93-9, ISIS 119849 404880-94-0, ISIS 119850 119848 404880-96-2, ISIS 119852 404880-97-3, ISIS 404880-95-1. ISIS 119851 404880-99-5, ISIS 119855 119853 404880-98-4, ISIS 119854 404881-00-1, ISIS 119856 404881-01-2, ISIS 119857 404881-02-3, ISIS 404881-04-5, ISIS 119860 119858 404881-03-4, ISIS 119859 404881-05-6, ISIS 119861 404881-06-7, ISIS 119862 404881-07-8, ISIS 404881-09-0, ISIS 119865 404881-08-9, ISIS 119864 119863 404881-12-5, ISIS 404881-10-3, ISIS 119866 404881-11-4, ISIS 119867 404881-13-6, ISIS 119869 404881-14-7, ISIS 119870 119868 404881-16-9, ISIS 119872 404881-17-0, ISIS 404881-15-8. ISIS 119871 404881-18-1, ISIS 119874 404881-19-2, ISIS 119875 119873 404881-20-5, ISIS 119876 404881-21-6. ISIS 119877 404881-23-8, ISIS 404881-24-9, ISIS 119880 404881-25-0, ISIS 119881 119879 404881-26-1, ISIS 119882 404881-27-2, ISIS 119883 404881-28-3. ISIS 404881-29-4, ISIS 119885 404881-30-7, ISIS 119886 119884 404881-31-8, ISIS 119887 404881-32-9, ISIS 119888 404881-33-0, ISIS 404881-35-2, ISIS 119891 404881-34-1, ISIS 119890 119889 404881-36-3, ISIS 119892 404881-37-4, ISIS 119893 404881-38-5, ISIS 404881-39-6, ISIS 119895 119894 404881-40-9, ISIS 119896 404881-41-0, ISIS 119897 404881-42-1, ISIS 119898 404881-43-2, ISIS 119899 404881-44-3, ISIS 119900 404881-45-4, ISIS 119901 404881-46-5, ISIS 119903 404881-47-6, ISIS 119904 404881-48-7, ISIS 119905 404881-49-8, ISIS 119906 404881-50-1, ISIS 119907 404881-51-2, ISIS 119908 404881-52-3, ISIS 119909 404881-53-4, ISIS 404881-55-6, ISIS 119912 404881-54-5, ISIS 119911 119910 404881-57-8, ISIS 119914 404881-58-9, ISIS 119915 404881-59-0, ISIS 404881-60-3, ISIS 119917 404997-71-3, ISIS 119992 119916 404997-75-7, ISIS 119921 404997-76-8, ISIS 404997-73-5, ISIS 119919 119922 404997-77-9, ISIS 119925 404997-78-0, ISIS 119926 404997-79-1, ISIS 119927 404997-80-4, ISIS 119928 404997-81-5, ISIS 119929 404997-82-6, ISIS 119930 404997-83-7, ISIS 119931 404997-84-8, ISIS 119932 404997-85-9, ISIS 119933 404997-86-0, ISIS 404997-87-1, ISIS 119935 404997-88-2, ISIS 119936 119934 404997-89-3, ISIS 119937 404997-90-6, ISIS 119938 404997-91-7, ISIS 404997-92-8, ISIS 119940 404997-93-9, ISIS 119941 119939 404997-95-1, ISIS 119943 404997-96-2, ISIS 404997-94-0, ISIS 119942 404997-98-4, ISIS 119946 119944 404997-97-3, ISIS 119945 404997-99-5, ISIS 119947 404998-00-1, ISIS 119948 404998-01-2, ISIS 404998-03-4, ISIS 119951 404998-04-5, ISIS 119952 119949 404998-05-6, ISIS 119953 404998-06-7, ISIS 119954 404998-08-9, ISIS 119956 404998-09-0, ISIS 119957 404998-10-3, ISIS 119958 404998-11-4, ISIS 119959 404998-12-5, ISIS 119960 404998-13-6, ISIS 404998-15-8, ISIS 119963 404998-16-9, ISIS 119964 404998-17-0, ISIS 119965 404998-18-1, ISIS 119966 404998-19-2, ISIS 404998-20-5, ISIS 119968 404998-21-6, ISIS 119969 119967 404998-23-8, ISIS 119971 404998-24-9, ISIS 404998-22-7, ISIS 119970 119972 404998-25-0, ISIS 119973 404998-26-1, ISIS 119974 404998-27-2, ISIS 119975 404998-28-3, ISIS 119976 404998-29-4, ISIS 119977 404998-30-7, ISIS 119978 404998-31-8, ISIS 119979 404998-33-0, ISIS 119981 404998-34-1, ISIS 404998-32-9, ISIS 119980 404998-36-3, ISIS 119984 404998-35-2, ISIS 119983 119982 404998-37-4, ISIS 119985 404998-39-6, ISIS 119987 404998-40-9, ISIS 404998-41-0, ISIS 119989 404998-42-1, ISIS 119990 404998-43-2, ISIS 119991 404998-44-3, ISIS 119993 404998-45-4, ISIS 404998-47-6, ISIS 119996 404998-46-5, ISIS 119995 119994 404998-49-8, ISIS 119998 404998-50-1, ISIS 404998-48-7, ISIS 119997 119999 RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (BH3 interacting domain death agonist-targeted; identification of genetic targets for modulation by antisense oligonucleotides and generation of oligonucleotides for gene modulation) 247116-73-0 247162-49-8 247162-51-2 247211-92-3 247211-94-5 TT 247212-33-5 247212-36-8 250225-18-4 634220-94-3. ISIS 247212-29-9 18624 634220-95-4, ISIS 19211 634220-96-5, ISIS 19213 634220-97-6, ISIS 19217 634220-98-7, ISIS 19219 634220-99-8, ISIS 19220 634221-00-4, ISIS 19221 634221-01-5, ISIS 19222 634221-02-0 634221-00-4, ISIS 19221 634221-02-6, ISIS 19223 634221-03-7, ISIS 19225 634221-04-8, ISIS 19230 634221-05-9. ISIS 19233 634221-06-0, ISIS 19235 634221-07-1, ISIS 19236

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        (CD40-targeted; identification of genetic targets for modulation by
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     RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (Jagged 2-targeted; identification of genetic targets for modulation by
        antisense oligonucleotides and generation of oligonucleotides for gene
       modulation)
     256631-53-5, ISIS 29576 256631-54-6, ISIS 29577 256631 29578 256631-56-8, ISIS 29579 256631-59-1, ISIS 29582
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            634221-39-9, ISIS 29612
     RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (PTEN phosphatase-targeted; identification of genetic targets for
        modulation by antisense oligonucleotides and generation of
        oligonucleotides for gene modulation)
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     147801 497213-53-3, ISIS 147806 497213-56-6, ISIS 147809
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     497213-72-6, ISIS 147825 497213-73-7, ISIS 147826 497213-76-0, ISIS
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     497213-86-2, ISIS 147480 497213-88-4, ISIS 147482
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     497214-13-8, ISIS 147507 497214-19-4, ISIS 147764
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    497214-30-9, ISIS 147775 497214-31-0, ISIS 147776
    RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (apolipoprotein B-targeted; identification of genetic targets for
       modulation by antisense oligonucleotides and generation of
       oligonucleotides for gene modulation)
    50-99-7, D-Glucose, analysis 57-88-5, Cholesterol, analysis 9013-08-5, Phosphoenolpyruvate carboxykinase
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    141907-41-7, Matrix metalloproteinase 186322-81-6, Caspase
    RL: ARU (Analytical role, unclassified); ANST (Analytical study)
        (assay system; identification of genetic targets for modulation by
       antisense oligonucleotides and generation of oligonucleotides for gene
       modulation)
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    148640-14-6, Protein kinase Akt-1
                                        260402-76-4, Protein kinase ELK1
    301166-54-1, Gene PTEN phosphatase
    RL: BSU (Biological study, unclassified); BIOL (Biological study) (target gene encoding; identification of genetic targets for modulation
       by antisense oligonucleotides and generation of oligonucleotides for
       gene modulation)
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    493667-07-5, ISIS 155659 493667-08-6, ISIS 155660 493667-09-7, ISIS
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    155700 493667-40-6, ISIS 155701 493667-41-7, ISIS 155702
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    RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (transforming growth factor .beta.3-targeted; identification of genetic
       targets for modulation by antisense oligonucleotides and generation of
       oligonucleotides for gene modulation)
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634242-68-5

634242-64-1

634242-69-6

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        (unclaimed nucleotide sequence; identification of genetic targets for
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634242-60-7

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     139:185625
     Entered STN: 15 Aug 2003
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    Little, Andrew S.; Lamparski, Henry G.; Henderson, Daniel R.; Schuur, Eric
    U.S. Pat. Appl. Publ., 115 pp., Cont.-in-part of U.S. Ser. No. 509,591.
     CODEN: USXXCO
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     ICM A61K048-00
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    The present invention provides compns. comprising host cell specific
    adenovirus vehicles for transfecting target host cells. The compns.
    comprise replication competent adenovirus having an adenovirus gene
    essential for replication under transcriptional control of a cell type
     specific transcriptional response element (TRE) and polyethylene glycol
     (PEG) as a masking agent.
    tissue specific replication competent adenovirus vector drug carrier
ST
IT
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        (E1A; compns. comprising tissue specific replication competent
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IT
    Gene, microbial
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (E1B; compns. comprising tissue specific replication competent
       adenoviral vectors)
TΤ
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       adenoviral vectors)
IT
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        (MUC1, gene, transcriptional regulatory element of; compns. comprising
        tissue specific replication competent adenoviral vectors)
    Genetic element
IT
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        (TRE (transcriptional response element); compns. comprising tissue
        specific replication competent adenoviral vectors)
TТ
    Drug delivery systems
        (carriers; compns. comprising tissue specific replication competent
       adenoviral vectors)
IT
    Prostate gland
        (cell specific TRE; compns. comprising tissue specific replication
       competent adenoviral vectors)
TT
    Adenoviral vectors
    DNA sequences
    Gene therapy
    Human
        (compns. comprising tissue specific replication competent adenoviral
       vectors)
IT
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IT
    Carcinoembryonic antigen
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        adenoviral vectors)
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    Animal cell
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        adenoviral vectors)
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     Enhancer (genetic element)
     Promoter (genetic element)
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        replication competent adenoviral vectors)
IT
     Proteins
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        (probasins, gene, transcriptional regulatory element of; compns. comprising tissue specific replication competent adenoviral vectors)
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        (vector; compns. comprising tissue specific replication competent
        adenoviral vectors)
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    9001-01-8, Kallikrein
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (2 gene, human, transcriptional regulatory element of; compns.
        comprising tissue specific replication competent adenoviral vectors)
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     79-09-4D, Propanoic acid, succinimidyl, biological studies 110-14-5D,
     Succinamide, succinimidyl 102696-21-9, Succinimidyl succinate RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (compns. comprising tissue specific replication competent adenoviral
        vectors)
TΤ
     25322-68-3, Polyethylene glycol
     RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (compns. comprising tissue specific replication competent adenoviral
        vectors)
     6066-82-6D, N-Hydroxysuccinimide, active esters
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CN
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*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
L27
     ANSWER 10 OF 25 HCAPLUS COPYRIGHT 2004 ACS on STN
     2003:590814 HCAPLUS
AN
     139:112801
DN
ED
     Entered STN:
                   01 Aug 2003
     Protein and cDNA sequences of a human cytokine CD40-ligand lacking
ΤI
     native-pattern glycosylation
     Armitage, Richard J.; Fanslow, William C.; Spriggs, Melanie K.
IN
PA
     USA
     U.S. Pat. Appl. Publ., 63 pp., Cont.-in-part of U.S. Ser. No. 392,618,
     abandoned.
     CODEN: USXXCO
DT
     Patent
LΑ
     English
IC
     ICM A61K038-17
     ICS C07K014-705; C12P021-02; C12N005-06; C07H021-04
     514008000; 530350000; 536023500; 435069100; 435320100; 435325000
     3-3 (Biochemical Genetics)
     Section cross-reference(s): 13, 15
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              GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
             LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM,
             PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN,
         TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
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PRAI US 1991-783707
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     US 1991-805723
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                                 19921023 <--
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     US 1999-392618
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     EP 1992-925017
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     US 2002-200242
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CLASS
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                         C07K014-705; C12P021-02; C12N005-06; C07H021-04
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                  NCL
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                         435325000
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                         C07K014/705Q; C12N015/62
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                         C07K014/705Q; C12N015/62
AB The invention provides protein and cDNA sequences of a human cytokine
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CD40-ligand lacking native-pattern glycosylation. More particularly, this
     invention provides isolated human and murine CD40-L polypeptides that bind
     to the extracellular binding region of a CD40 receptor and lack
    native-pattern glycosylation.
    sequence human cytokine CD40 ligand glycosylation
ST
     Cytokines
IT
    RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     PRP (Properties); BIOL (Biological study); PREP (Preparation)
        (CD40-ligand; protein and cDNA sequences of human cytokine CD40-ligand
        lacking native-pattern glycosylation)
    Gene, animal
    RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
     (Biological study)
        (for CD40-ligand; protein and cDNA sequences of human cytokine
        CD40-ligand lacking native-pattern glycosylation)
    Protein motifs
TT
        (leucine zipper; protein and cDNA sequences of human cytokine
        CD40-ligand lacking native-pattern glycosylation)
IT
    Glycosylation
    Human
    Molecular cloning
    Mutation
     Protein sequences
     cDNA sequences
        (protein and cDNA sequences of human cytokine CD40-ligand lacking
        native-pattern glycosylation)
IT
    CD40 (antigen)
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (protein and cDNA sequences of human cytokine CD40-ligand lacking
        native-pattern glycosylation)
     562120-99-4P, Cytokine CD40-ligand (human)
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     PRP (Properties); BIOL (Biological study); PREP (Preparation)
        (amino acid sequence; protein and cDNA sequences of human cytokine
        CD40-ligand lacking native-pattern glycosylation)
     562121-48-6
     RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
     (Biological study)
        (leucine zipper domain sequence; protein and cDNA sequences of human
        cytokine CD40-ligand lacking native-pattern glycosylation)
     562120-98-3
     RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
     (Biological study)
        (nucleotide sequence; protein and cDNA sequences of human cytokine
        CD40-ligand lacking native-pattern glycosylation)
                  562122-85-4
                                 562122-86-5 562122-87-6 562122-88-7
IT
     562122-83-2
                   562122-90-1
     562122-89-8
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     562122-94-5
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                                                             562123-00-6
     RL: PRP (Properties)
        (unclaimed nucleotide sequence; protein and cDNA sequences of a human
        cytokine CD40-ligand lacking native-pattern glycosylation)
                  562122-95-6 562122-99-0
IT
     562122-84-3
                                               562123-01-7
     RL: PRP (Properties)
        (unclaimed protein sequence; protein and cDNA sequences of a human
        cytokine CD40-ligand lacking native-pattern glycosylation)
                               245427-08-1
     98849-88-8 203244-38-6
IT
     RL: PRP (Properties)
        (unclaimed sequence; protein and cDNA sequences of a human cytokine
        CD40-ligand lacking native-pattern glycosylation)
IT
     562122-88-7
    RL: PRP (Properties)
        (unclaimed nucleotide sequence; protein and cDNA sequences of a human
        cytokine CD40-ligand lacking native-pattern glycosylation)
     562122-88-7 HCAPLUS
RN
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CN
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*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
L27 ANSWER 11 OF 25 HCAPLUS COPYRIGHT 2004 ACS on STN
     2003:545709 HCAPLUS
    139:79122
DN
ED
     Entered STN: 17 Jul 2003
    A new calicivirus "San Miguel sea lion virus 5 serotype Homosapien-1"
TI
     (SMSV-5 Hom-1), and diagnosis, prevention and treatment of calicivirus
     infection in humans
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Smith, Alvin W.
IN
     The State of Oregon Acting by and Through the State Board of Higher
     Education On Behalf of Oregon State University, USA
     U.S., 48 pp.
SO
     CODEN: USXXAM
ידת
     Patent
     English
LA
     ICM C12Q001-70
IC
     ICS G01N033-53; C07K017-00
NCL
     435005000; 435007100; 530350000
     1-5 (Pharmacology)
     Section cross-reference(s): 10
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                                                                      DATE
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US 6593080
                         C120001-70
                         G01N033-53; C07K017-00
                 ICS
                         435005000; 435007100; 530350000
                 NCL
    Caliciviruses are herein recognized as potentially important human
     pathogens, infecting a large percentage (18%) of a sampled human
     population. A new calicivirus isolate called "San Miguel sea lion virus 5
     serotype Homosapien-1" (SMSV-5 Hom-1) is also disclosed. Compns. and
     methods are disclosed that allow diagnosis, prevention and treatment of
     calicivirus (including SMSV-5 Hom-1) infection in humans.
ST
     calicivirus SMSV5Hom1 human infection diagnosis
IT
     Calicivirus
        (SMSV-5 Hom-1; new calicivirus "San Miguel sea lion virus 5 serotype Homosapien-1" (SMSV-5 Hom-1), and diagnosis, prevention and treatment
        of calicivirus infection in humans)
IT
     Thymus gland, disease
        (atrophy, detecting calicivirus in samples from; new calicivirus "San
        Miguel sea lion virus 5 serotype Homosapien-1" (SMSV-5 Hom-1), and
        diagnosis, prevention and treatment of calicivirus infection in humans)
IT
        (calicivirus; new calicivirus "San Miguel sea lion virus 5 serotype Homosapien-1" (SMSV-5 Hom-1), and diagnosis, prevention and treatment
        of calicivirus infection in humans)
     Fatique, biological
IT
        (chronic fatigue syndrome, detecting calicivirus in samples from; new
        calicivirus "San Miguel sea lion virus 5 serotype Homosapien-1" (SMSV-5
        \mbox{{\tt Hom-1}}\mbox{), and diagnosis, prevention and treatment of calicivirus}
        infection in humans)
IT
     Blister
     Diarrhea
     Encephalitis
     Hemorrhage
     Hepatitis
     Pneumonia
        (detecting calicivirus in samples from; new calicivirus "San Miquel sea
        lion virus 5 serotype Homosapien-1" (SMSV-5 Hom-1), and diagnosis,
        prevention and treatment of calicivirus infection in humans)
IT
     Mucous membrane
        (disease, erosion, detecting calicivirus in samples from; new
        calicivirus "San Miguel sea lion virus 5 serotype Homosapien-1" (SMSV-5
        Hom-1), and diagnosis, prevention and treatment of calicivirus
        infection in humans)
IT
     Proteins
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     PRP (Properties); BIOL (Biological study); PREP (Preparation)
        (encoded by p5RT73 genomic segment; new calicivirus "San Miguel sea
lion virus 5 serotype Homosapien-1" (SMSV-5 Hom-1), and diagnosis,
        prevention and treatment of calicivirus infection in humans)
IT
     Diagnosis
        (mol.; new calicivirus "San Miguel sea lion virus 5 serotype
        Homosapien-1" (SMSV-5 Hom-1), and diagnosis, prevention and treatment
        of calicivirus infection in humans)
IT
     Heart, disease
        (myocarditis, detecting calicivirus in samples from; new calicivirus
        "San Miguel sea lion virus 5 serotype Homosapien-1" (SMSV-5 Hom-1), and
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diagnosis, prevention and treatment of calicivirus infection in humans)
IT
     Muscle, disease
         (myositis, detecting calicivirus in samples from; new calicivirus "San
        Miguel sea lion virus 5 serotype Homosapien-1" (SMSV-5 Hom-1), and
         diagnosis, prevention and treatment of calicivirus infection in humans)
IT
     Antiviral agents
     Human
     Immunostimulants
     Molecular cloning
     Protein'sequences
         (new calicivirus "San Miguel sea lion virus 5 serotype Homosapien-1"
         (SMSV-5 Hom-1), and diagnosis, prevention and treatment of calicivirus
         infection in humans)
IT
     Antigens
     RL: ARG (Analytical reagent use); BSU (Biological study, unclassified);
     DGN (Diagnostic use); PRP (Properties); THU (Therapeutic use); ANST
     (Analytical study); BIOL (Biological study); USES (Uses)
         (new calicivirus "San Miguel sea lion virus 5 serotype Homosapien-1"
         (SMSV-5 Hom-1), and diagnosis, prevention and treatment of calicivirus
         infection in humans)
ΙT
     Gene, microbial
     RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
     (Biological study)
         (p5RT73 genomic segment; new calicivirus "San Miguel sea lion virus 5
        serotype Homosapien-1" (SMSV-5 Hom-1), and diagnosis, prevention and treatment of calicivirus infection in humans)
IT
     Abortion
         (spontaneous, detecting calicivirus in samples from; new calicivirus
         "San Miguel sea lion virus 5 serotype Homosapien-1" (SMSV-5 Hom-1), and
         diagnosis, prevention and treatment of calicivirus infection in humans)
IT
     Antibodies and Immunoglobulins
     RL: ARG (Analytical reagent use); DGN (Diagnostic use); ANST (Analytical
     study); BIOL (Biological study); USES (Uses)
        (to calicivirus; new calicivirus "San Miguel sea lion virus 5 serotype Homosapien-1" (SMSV-5 Hom-1), and diagnosis, prevention and treatment
         of calicivirus infection in humans)
IT
     Organelle
         (vesicle, detecting calicivirus in samples from; new calicivirus "San
         Miguel sea lion virus 5 serotype Homosapien-1" (SMSV-5 Hom-1), and
        diagnosis, prevention and treatment of calicivirus infection in humans)
IT
        (viral, diagnosis and treatment of; new calicivirus "San Miguel sea
lion virus 5 serotype Homosapien-1" (SMSV-5 Hom-1), and diagnosis,
        prevention and treatment of calicivirus infection in humans)
IT
     556161-83-2P, Antigen (calicivirus)
                                               556162-56-2P
                                                               556162-57-3P
     556162-58-4P
     RL: ARG (Analytical reagent use); BPN (Biosynthetic preparation); BSU (Biological study, unclassified); PRP (Properties); ANST (Analytical
     study); BIOL (Biological study); PREP (Preparation); USES (Uses)
        (amino acid sequence; new calicivirus "San Miguel sea lion virus 5 serotype Homosapien-1" (SMSV-5 Hom-1), and diagnosis, prevention and
         treatment of calicivirus infection in humans)
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     US6593080 SEQID: 3 unclaimed DNA 556164-58-0, 4: PN: US6593080 SEQID: 4
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     556164-62-6 556165-87-8
     RL: PRP (Properties)
         (unclaimed nucleotide sequence; new calicivirus "San Miguel sea lion
        virus 5 serotype Homosapien-1" (SMSV-5 Hom-1), and diagnosis,
        prevention and treatment of calicivirus infection in humans)
IT
     556164-56-8 556165-88-9 556165-89-0
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     556165-92-5
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     RL: PRP (Properties)
         (unclaimed protein sequence; new calicivirus "San Miguel sea lion virus
        5 serotype Homosapien-1" (SMSV-5 Hom-1), and diagnosis, prevention and
        treatment of calicivirus infection in humans)
IT
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         (unclaimed sequence; new calicivirus "San Miguel sea lion virus 5
        serotype Homosapien-1" (SMSV-5 Hom-1), and diagnosis, prevention and
        treatment of calicivirus infection in humans)
RE.CNT 14
               THERE ARE 14 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE
(1) Berry; ASM Annual Meeting 1987
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(2) Berry; Abstract, 1987 ASM Annual Meeting 1987
(3) Bradley; Journal of Hepatology 1995, V22 (Suppl 1), P140
(4) Nakata; Journal of Clinical Microbiology 1983, V17, P198 MEDLINE
(5) Neill; Genbank Accession No P89669 1997
(6) Neill; Virus Research 1992, V24, P211 HCAPLUS (7) Neill; Virus Research 1992, V24, P211 HCAPLUS
(8) Poet, S; Thesis presented 1994
(9) Smith; Amer J Vet Res 1977, V38(1), P101 MEDLINE
(10) Smith; Amer J Vet Res 1978, V39(2), P287 MEDLINE
(11) Smith; Calicivirus Isolation and Persistence in a Pygmy Chimpanzee (Pan
    paniscus), 1983, V221, P79 MEDLINE
(12) Smith; Clinical Infectious Diseases 1998, V26, P424
(13) Smith; Emerging Infectious Diseases 1998, V4(1), P13 MEDLINE
(14) Soergel; Archives of Virology 1978, V57(3) HCAPLUS
     556164-59-1
     RL: PRP (Properties)
        (unclaimed nucleotide sequence; new calicivirus "San Miguel sea lion virus 5 serotype Homosapien-1" (SMSV-5 Hom-1), and diagnosis,
        prevention and treatment of calicivirus infection in humans)
RN
     556164-59-1 HCAPLUS
     CN
     (CA INDEX NAME)
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L27 ANSWER 12 OF 25 HCAPLUS COPYRIGHT 2004 ACS on STN
AN
     2003:260540 HCAPLUS
     138:249945
ED
     Entered STN: 04 Apr 2003
     Biallelic markers for use in constructing a high density disequilibrium
ΤI
     map of the human genome and for diagnosis of the risk for developing
     Alzheimer's disease
     Cohen, Daniel; Chumakov, Ilya; Blumenfeld, Marta
IN
PA
     Genset, Fr.
     U.S., 134 pp., Cont.-in-part of U.S. Ser. No. 298,850, abandoned.
SO
     CODEN: USXXAM
DT
     Patent
     English
LA
     C07H021-04; C07Q001-68
IC
     435006000; 435091200; 536023100; 536024300; 536023400
NCL
     3-3 (Biochemical Genetics)
     Section cross-reference(s): 13, 14
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             MD, RU, TJ, TM
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PRAI US 1998-82614P
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CT-ASS
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                                           C07Q001-68
 US 6537751
                         435006000; 435091200; 536023100; 536024300; 536023400
                 NCT.
                 ECLA C12Q001/68B6; C12Q001/68B10A; C12Q001/68M
 WO 9954500
    The present invention relates to genomic maps comprising biallelic
     markers, new biallelic markers, and methods of using biallelic markers.
     Primers hybridizing to regions flanking these biallelic markers are also
     provided. This invention provides polynucleotides and methods suitable
     for genotyping a nucleic acid containing sample for one or more biallelic
     markers of the invention. Further, the invention provides a number of
     methods utilizing the biallelic markers of the invention including methods
     to detect a statistical correlation between a biallelic marker allele and
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a phenotype and/or between a biallelic marker haplotype and a phenotype.

Nguyen 10/068160 The compns. and methods of the invention also find use in the identification of targets for the development of pharmaceutical agents and diagnostic methods, as well as the characterization of differential efficacious responses to and side effects from pharmaceutical agents acting on a disease (e.g., Alzheimer's disease, prostate cancer, or asthma) as well as other treatments. The invention provides 3934 biallelic marker sequences, as well as the primer pairs for amplification and detection of each marker. In particular, an allele at position 24 of biallelic marker 99-365-344 is associated with an increased risk for developing Alzheimer's disease. [This abstract record is one of three records for this document necessitated by the large number of index entries required to fully index the document and publication system constraints.]. biallelic marker human genome mapping; sequence biallelic marker DNA human genome; PCR primer biallelic marker human genome; drug Alzheimers asthma gene biallelic marker Gene, animal RL: ANT (Analyte); DGN (Diagnostic use); ANST (Analytical study); BIOL (Biological study); USES (Uses) (APOE, determination of haplotype frequency; biallelic markers for use in constructing a high d. disequil. map of the human genome and for diagnosis of the risk for developing Alzheimer's disease) Computer application DNA sequences Drug design Drug screening Genetic mapping Genetic markers Genome Genotyping (method) Human Nucleic acid amplification (method) Nucleic acid hybridization PCR (polymerase chain reaction) (biallelic markers for use in constructing a high d. disequil. map of the human genome and for diagnosis of the risk for developing Alzheimer's disease) Primers (nucleic acid) RL: ARG (Analytical reagent use); DGN (Diagnostic use); ANST (Analytical study); BIOL (Biological study); USES (Uses) (biallelic markers for use in constructing a high d. disequil. map of the human genome and for diagnosis of the risk for developing Alzheimer's disease) Anti-Alzheimer's agents Antiasthmatics (design and screening of; biallelic markers for use in constructing a high d. disequil. map of the human genome and for diagnosis of the risk for developing Alzheimer's disease) Allele frequency (determination in a population; biallelic markers for use in constructing a high d. disequil. map of the human genome and for diagnosis of the risk for developing Alzheimer's disease) Haplotypes (determination of frequency of; biallelic markers for use in constructing a high d. disequil. map of the human genome and for diagnosis of the risk for developing Alzheimer's disease) Test kits (diagnostic; biallelic markers for use in constructing a high d. disequil. map of the human genome and for diagnosis of the risk for developing Alzheimer's disease) Genetic linkage (disequil.; biallelic markers for use in constructing a high d. disequil. map of the human genome and for diagnosis of the risk for developing Alzheimer's disease) Genetic methods (enzyme-based mismatch detection assay; biallelic markers for use in constructing a high d. disequil. map of the human genome and for diagnosis of the risk for developing Alzheimer's disease) (genetic; biallelic markers for use in constructing a high d. disequil. map of the human genome and for diagnosis of the risk for developing Alzheimer's disease) Prostate gland, neoplasm (inhibitors, design and screening of; biallelic markers for use in constructing a high d. disequil. map of the human genome and for

IT

TΨ

IT

IT

TT

IT

IT

IT

IT

IT

Antitumor agents

diagnosis of the risk for developing Alzheimer's disease)

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(prostate gland, design and screening of; biallelic markers for use in
        constructing a high d. disequil. map of the human genome and for
        diagnosis of the risk for developing Alzheimer's disease)
TТ
     Alzheimer's disease
        (screening for; biallelic markers for use in constructing a high d.
        disequil. map of the human genome and for diagnosis of the risk for
        developing Alzheimer's disease)
IT
     Genomic library
     cDNA library
        (screening of; biallelic markers for use in constructing a high d.
        disequil. map of the human genome and for diagnosis of the risk for
        developing Alzheimer's disease)
TT
     Genetic polymorphism
        (single nucleotide; biallelic markers for use in constructing a high d.
        disequil. map of the human genome and for diagnosis of the risk for
        developing Alzheimer's disease)
                   502238-05-3
IT
                                 502238-06-4
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     (Analytical study); BIOL (Biological study); USES (Uses)
        (downstream amplification primer for biallelic marker; biallelic
        markers for use in constructing a high d. disequil. map of the human
        genome and for diagnosis of the risk for developing Alzheimer's
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(Analytical study); BIOL (Biological study); USES (Uses)
   (downstream amplification primer for biallelic marker; biallelic
   markers for use in constructing a high d. disequil. map of the human
   genome and for diagnosis of the risk for developing Alzheimer's
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     (Analytical study); BIOL (Biological study); USES (Uses)
        (downstream amplification primer for biallelic marker; biallelic
        markers for use in constructing a high d. disequil. map of the human
        genome and for diagnosis of the risk for developing Alzheimer's
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        (downstream amplification primer for biallelic marker; biallelic
        markers for use in constructing a high d. disequil. map of the human
        genome and for diagnosis of the risk for developing Alzheimer's
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        (downstream amplification primer for biallelic marker; biallelic
        markers for use in constructing a high d. disequil. map of the human
        genome and for diagnosis of the risk for developing Alzheimer's
        disease)
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        (downstream amplification primer for biallelic marker; biallelic
        markers for use in constructing a high d. disequil. map of the human
        genome and for diagnosis of the risk for developing Alzheimer's
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        (downstream amplification primer for biallelic marker; biallelic
        markers for use in constructing a high d. disequil. map of the human
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        (downstream amplification primer for biallelic marker; biallelic
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TТ
    Tissue specific adenoviral vectors
    Henderson, Daniel R.; Schuur, Eric R.
IN
PA
    Cell Genesys, Inc., USA
    U.S. Pat. Appl. Publ., 83 pp., Cont.-in-part of U.S. Ser. No. 33,555,
SO
    abandoned.
     CODEN: USXXCO
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    English
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NCL
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C07K014/075; C07K014/47A6; C12N015/63; C12N015/86F;
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US 2002164799
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    Host cell specific adenovirus vehicles are provided for transfecting
     target host cells. By providing for transcriptional initiating regulation
     dependent upon transcription factors that are only active in specific,
     limited cell types, virus replication will be restricted to the target
     cells. The modified adenovirus may be used as a vehicle for introducing
    new genetic capability, particularly associated with cytotoxicity for
    treating neoplasia.
ST
     genetic engineering gene therapy adenoviral vector tissue targeting
    Mammary gland, neoplasm
     Prostate gland
        (-specific transcriptional response element; tissue-specific adenoviral
        vectors)
     Carcinoembryonic antigen
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    RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (E1B; tissue-specific adenoviral vectors)
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        (E4; tissue-specific adenoviral vectors)
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        (TRE (transcriptional response element); tissue-specific adenoviral
        vectors)
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    Intestine, neoplasm
        (colon, -specific transcriptional response element; tissue-specific
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    Protein motifs
        (death domain; tissue-specific adenoviral vectors)
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        (late; tissue-specific adenoviral vectors)
    Adenoviral vectors
     Antitumor agents
    Gene targeting
     Gene therapy
     Genetic engineering
    Transformation, genetic
        (tissue-specific adenoviral vectors)
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    Antisense oligonucleotides inhibiting aldehyde dehydrogenase gene
ΤI
     expression for use in treatment of alcohol abuse
    Garver, Eric; Tu, Guang-chou; Israel, Yedy
TN
PA
     Thomas Jefferson University, USA
so
    U.S. Pat. Appl. Publ., 66 pp., Cont.-in-part of U.S. 6,277,981.
    CODEN: USXXCO
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    536023200; 514044000
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                ECLA
                       C12N015/11B; C12N015/11B3
    The invention includes methods for reducing alc. consumption in mammals by
AB
     inhibiting expression of an aldehyde dehydrogenase (e.g., liver
     mitochondrial aldehyde dehydrogenase) in cells of the mammal. Expression
     of the aldehyde dehydrogenase can be inhibited by administering (locally
     or systemically) an antisense oligonucleotide (e.g., one 12-2000 residues
     in length) to the mammal, whereby expression of a relatively active allele
     of an aldehyde dehydrogenase gene can be inhibited. Aldehyde
     dehydrogenase activity can also be inhibited in the cells of a mammal, for
     the same purposes, by administering to the cells an expression vector
     encoding an inactive allele dominant of the aldehyde dehydrogenase. Upon
     expression, subunits of the protein encoded by the inactive allele can
     coalesce with one another or with subunits of the cells normally-expressed
     aldehyde dehydrogenase to lower the level of aldehyde dehydrogenase
     activity in the cell. The invention also relates to methods for
     predicting whether an antisense oligonucleotide (ASO) will be efficacious
     for inhibiting expression of a gene. Thus, oligonucleotides targeting
     GGGA motifs seem to be particularly effective.
     antisense oligonucleotide efficacy; alc abuse aldehyde dehydrogenase ALDH2
ST
     gene antisense oligonucleotide
    Gene, animal
IT
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (ALDH2-1; antisense oligonucleotides inhibiting aldehyde dehydrogenase
        gene expression for use in treatment of alc. abuse)
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    Gene, animal
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (ALDH2-2, enzyme inhibition by expression of; antisense
        oligonucleotides inhibiting aldehyde dehydrogenase gene expression for
        use in treatment of alc. abuse)
TT
    Gene, animal
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (ALDH2; antisense oligonucleotides inhibiting aldehyde dehydrogenase
        gene expression for use in treatment of alc. abuse)
IT
    Alcoholism
    Human
        (antisense oligonucleotides inhibiting aldehyde dehydrogenase gene
        expression for use in treatment of alc. abuse)
    Antisense oligonucleotides
TΤ
     RL: PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES
     (Uses)
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(antisense oligonucleotides inhibiting aldehyde dehydrogenase gene
        expression for use in treatment of alc. abuse)
IT
     Tumor necrosis factors
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (antisense oligonucleotides inhibiting tumor necrosis factor .alpha.
        gene expression)
IT
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
(expression; method for producing effective antisense oligonucleotides
        for inhibition of gene expression)
TT
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        gene expression)
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AN
    137:334622
DN
ED
    Entered STN: 31 Oct 2002
    Engineering of cyclodextrin glycosyltransferase deletion mutants for
    improved production of gamma cyclodextrin
    Schulz, Georg E.; Parsiegla, Goetz
IN
    Consortium Fuer Elektrochemische Industrie G.m.b.H., Germany
PA
    U.S., 24 pp., Cont.-in-part of U.S. Ser. No. 816,317. CODEN: USXXAM
so
DT
    Patent
LΑ
    English
    ICM C12N009-10
IC
        C12N001-00; C12P021-04; C08H001-00; C07H021-04
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CC
    7-2 (Enzymes)
     Section cross-reference(s): 3, 10
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                NCL
                       435193000
US 6472192
                ECLA
                       C12N009/10D1A19
                ECLA
                      C12N009/10D1A19
DE 19615336
    CGTases, which, when converting starch or starch-like substrates to CD,
    produce .gamma.-CD to an increased extent and still exhibit at least 60%
     of the specific total CGTase activity of the starting CGTase which was
    used for preparing the enzyme concerned. The amino acid sequences differ
     from the amino acid sequences of known CGTases by the deletion of from 3
     to 8 amino acids from the region from amino acid position 155 up to and
     including amino acid position 195, where position 1 of the protein
     sequence is the beginning of the signal peptide of the CGTase and the
     deletion brings about the increase in the .gamma .- CGTase activity of the
     protein. In particular, cyclodextrin glycosyltransferase deletion
     variants of Bacillus sp.1-1, Bacillus circulans, Bacillus ohbensis,
     Paenibacillus macerans and Thermanerobacter are described.
    qlycosyltransferase cyclodextrin deletion mutant recombinant; cyclodextrin
ST
     gamma starch recombinant glycosyltransferase mutant; sequence cyclodextrin
    glycosyltransferase Bacillus Thermoanerobacter Paenibacillus
     Bacillus circulans
    Bacillus ohbensis
     Paenibacillus macerans
     Thermoanaerobacter
        (CGTase of; engineering of cyclodextrin glycosyltransferase deletion
       mutants for improved production of gamma cyclodextrin)
IT
    Microorganism
        (CGTase; engineering of cyclodextrin glycosyltransferase deletion
       mutants for improved production of gamma cyclodextrin)
IT
     Peptides, biological studies
     RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
     (Biological study)
        (cyclodextrin glycosyltransferase; engineering of cyclodextrin
        glycosyltransferase deletion mutants for improved production of gamma
        cyclodextrin)
ΤT
    Mutation
        (deletion; engineering of cyclodextrin glycosyltransferase deletion
        mutants for improved production of gamma cyclodextrin)
IT
    Molecular cloning
        (engineering of cyclodextrin glycosyltransferase deletion mutants for
        improved production of gamma cyclodextrin)
TT
    Gene, microbial
    RL: BUU (Biological use, unclassified); PRP (Properties); BIOL (Biological
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        (for cyclodextrin glycosyltransferase; engineering of cyclodextrin
        glycosyltransferase deletion mutants for improved production of gamma
        cyclodextrin)
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Protein engineering
IT
         (of cyclodextrin glycosyltransferase deletion variants; engineering of
         cyclodextrin glycosyltransferase deletion mutants for improved production
        of gamma cyclodextrin)
     Protein sequences
IT
         (of cyclodextrin glycosyltransferase variants of Bacillus,
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         cyclodextrin glycosyltransferase deletion mutants for improved production
        of gamma cyclodextrin)
IT
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         (of cyclodextrin glycosyltransferase; engineering of cyclodextrin
        glycosyltransferase deletion mutants for improved production of gamma
        cyclodextrin)
IT
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         (sp. 1-1, CGTase of; engineering of cyclodextrin glycosyltransferase
         deletion mutants for improved production of gamma cyclodextrin)
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     (Preparation); USES (Uses)
        (amino acid sequence; engineering of cyclodextrin glycosyltransferase deletion mutants for improved production of gamma cyclodextrin)
IT
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         (as substrate for cyclodextrin glycosyltransferase; engineering of
         cyclodextrin glycosyltransferase deletion mutants for improved production
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12619-70-4P, Cyclodextrin 17465-86-0P, .gamma.-Cyclodextrin
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     (Preparation)
         (engineering of cyclodextrin glycosyltransferase deletion mutants for
         improved production of gamma cyclodextrin)
     9030-09-5DP, Cyclodextrin glycosyltransferase, deletion mutants
TT
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     CAT (Catalyst use); PRP (Properties); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
         (engineering of cyclodextrin glycosyltransferase deletion mutants for
         improved production of gamma cyclodextrin)
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        (unclaimed sequence; engineering of cyclodextrin glycosyltransferase deletion mutants for improved production of gamma cyclodextrin)
RE.CNT 28
               THERE ARE 28 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE
(1) Anon; EP 0291067 1988 HCAPLUS
(2) Anon; DE 4009822 1991 HCAPLUS
(3) Anon; WO 9114770 1991 HCAPLUS
(4) Anon; EP 0481513 1992
(5) Anon; EP 0614571 1994
(6) Anon; EP 0630567 1994
(7) Anon; DE 4324650 1995 HCAPLUS
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    Biology/Reagent System P138
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(16) Leung; Technique 1989, V1(1), P11
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(22) Nitschke; Appl Microgiol Biotechnol 1990, V33, P542 HCAPLUS
(23) Penninga; Biochemistry 1995, V34(10), P3368 HCAPLUS
(24) Sanger; Proc Natl Acad Sci USA P5463
(25) Schmid; US 4822874 A 1989 HCAPLUS
(26) Schmid; US 5409824 A 1995 HCAPLUS
(27) Schulz; US 5474917 A 1995 HCAPLUS
(28) Tomita; Journal of Fermentation and Bioengineering 1990, V70(3), P190
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         glycosyltransferase deletion mutants for improved production of gamma
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L27 ANSWER 16 OF 25 HCAPLUS COPYRIGHT 2004 ACS on STN
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     2002:669668 HCAPLUS
     137:210920
     Entered STN: 05 Sep 2002
ED
     Antisense oligonucleotides inhibiting human telomerase reverse
TΤ
     transcriptase in diagnosis and treatment of cancer
ΙN
     Cech, Thomas R.; Lingner, Joachim; Nakamura, Toru; Chapman, Karen B.;
     Morin, Gregg B.; Harley, Calvin B.; Andrews, William H.
     Geron Corporation, USA
PA
     U.S., 34 pp., Cont.-in-part of U.S. 6,166,178.
SO
     CODEN: USXXAM
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     English
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TC
     A01N043-04; A61K031-70; C02H021-04
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     1-6 (Pharmacology)
     Section cross-reference(s): 3, 7, 14, 63
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WO 9950279
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    The present invention provides TRT antisense oligonucleotides, methods of
     detecting TRT, methods of diagnosing telomerase-related conditions,
    methods of diagnosing and providing a prognosis for cancer, and methods of
     treating telomerase-related conditions, including cancer.
     telomerase reverse transcriptase antisense oligonucleotide diagnosis
ST
     treatment cancer; human telomerase reverse transcriptase cDNA sequence
IT
    Antitumor agents
    Human
    Neoplasm
    Nucleic acid hybridization
        (antisense oligonucleotides inhibiting human telomerase reverse
        transcriptase in diagnosis and treatment of cancer)
    DNA
IT
    RNA
     RL: ANT (Analyte); BSU (Biological study, unclassified); ANST (Analytical
     study); BIOL (Biological study)
        (antisense oligonucleotides inhibiting human telomerase reverse
        transcriptase in diagnosis and treatment of cancer)
IT
    Antisense oligonucleotides
     RL: DGN (Diagnostic use); PRP (Properties); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (antisense oligonucleotides inhibiting human telomerase reverse
        transcriptase in diagnosis and treatment of cancer)
IT
     Phosphorothioate oligonucleotides
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (antisense oligonucleotides; antisense oligonucleotides inhibiting
       human telomerase reverse transcriptase in diagnosis and treatment of
        cancer)
IT
    Diagnosis
        (cancer; antisense oligonucleotides inhibiting human telomerase reverse
        transcriptase in diagnosis and treatment of cancer)
IT
    cDNA sequences
        (for telomerase reverse transcriptase of human; antisense
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Search done by Noble Jarrell

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oligonucleotides inhibiting human telomerase reverse transcriptase in
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IT
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     RL: BSU (Biological study, unclassified); DGN (Diagnostic use); PRP
     (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (for telomerase reverse transcriptase; antisense oligonucleotides
        inhibiting human telomerase reverse transcriptase in diagnosis and
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IT
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     Prognosis
IT
        (of cancer, TRT antisense DNA in; antisense oligonucleotides inhibiting
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TТ
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        (telomerase reverse transcriptase; antisense oligonucleotides
        inhibiting human telomerase reverse transcriptase in diagnosis and
        treatment of cancer)
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RE
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(13) Villeponteau; US 5583016 A 1996 HCAPLUS
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    Coupled amplification and sequencing method for detection of disease
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IN
    Leushner, James; Hui, May; Dunn, James M.; Lacroix, Jean-Michel
    Visible Genetics Inc., Can.
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    U.S., 62 pp., Cont.-in-part of U.S. Ser. No. 9,483.
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    Evaluation of a sample for the presence and qual. nature of a
     microorganism can be performed in a single vessel by combining a natural
     abundance DNA sample with a sequencing mixture containing a primer pair, a
     thermally stable polymerase such as ThermoSequenaseTM which incorporates
     dideoxynucleotides into an extending nucleic acid polymer at a rate which
     is no less than about 0.4 times the rate of incorporation of
     deoxynucleotides, nucleotide triphosphate feedstocks, and a chain
    terminating nucleotide triphosphate. The mixture is processed through multiple thermal cycles for annealing, extension and denaturation to
     produce a product mixture which is analyzed by electrophoresis. The present
     invention also provides a composition comprising a mixture of four
     deoxynucleotide triphosphates and at least one dideoxynucleotide
     triphosphate corresponding to one of the four deoxynucleotide
     triphosphates in a mole ratio from 1:50 to 1:500. Also provided is a kit
     for detection of a target microorganism comprising, in packaged
     combination, the above composition, thermostable polymerase enzyme and a pair
    of primers.
    PCR microorganism diagnosis kit primer sequence
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    Nucleotides, biological studies
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        (DNA, fluorescent labeled; coupled amplification and sequencing method
        for detection of disease causing microorganism)
     Chlamydia trachomatis
    Human immunodeficiency virus 1
    Human papillomavirus
    Test kits
        (coupled amplification and sequencing method for detection of disease
        causing microorganism)
    Nucleic acid amplification (method)
IT
        (coupled amplification and sequencing; coupled amplification and
        sequencing method for detection of disease causing microorganism)
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    Human immunodeficiency virus
        (detection; coupled amplification and sequencing method for detection
        of disease causing microorganism)
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        (in the kit; coupled amplification and sequencing method for detection
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        (mol., of disease causing microorganism; coupled amplification and
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(2) Anon; WO 8907149 1989 HCAPLUS
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(39) Wiemann; Anal Biochem 1995, V224, P117 HCAPLUS
(40) Wiemann; Anal Biochem 1996, V234, P166 HCAPLUS
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     Nucleic acid amplification using modular branched primers
TI
     Ulanovsky, Levy; Raja, Mugasimangalam C.
IN
     U.S., 33 pp., Cont.-in-part of U.S. 5,627,037. CODEN: USXXAM
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    Methods and compns. which expand the options for making primers for use in
     amplifying nucleic acid segments are disclosed. The invention eliminates
     the step of custom synthesis of primers for Polymerase Chain Reactions
     (PCR). Instead of being custom-synthesized, a primer is replaced by a
     combination of two or more oligonucleotide modules selected from a
     pre-synthesized library. A modular combination of just two
     oligonucleotides can mimic the performance of a conventional, custom-made
     primer by matching the sequence of the priming site in the template. Each
     oligonucleotide has a segment (I) that matches one of the stretches within
     the priming site as well as another segment (II) which is complementary to
     a segment of another oligonucleotide. A primer may be formed by two
     oligonucleotides with complementary II segments, while the I segments of
     the two oligonucleotides combine to form the target-complementary primer.
     The resulting branched primers comprise a double-stranded stem portion and
     a single-stranded, target complementary portion. The branched primer
     forms a three-way junction when annealed to the target. Two branched
     primers, or a branched and a traditional primer, may be used for the
     amplification.
     PCR modular branched primer
ST
     Nucleic acid amplification (method)
     PCR (polymerase chain reaction)
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(nucleic acid amplification using modular branched primers)
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      328603-42-5
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      RL: PRP (Properties)
         (unclaimed nucleotide sequence; nucleic acid amplification using
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             THERE ARE 27 CITED REFERENCES AVAILABLE FOR THIS RECORD
 RE.CNT 27
 RE
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      NAME)
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      ANSWER 19 OF 25 HCAPLUS COPYRIGHT 2004 ACS on STN
L27
AN
      2000:736280 HCAPLUS
DN
      133:291951
 ED
      Entered STN: 18 Oct 2000
 TI
      Cloning and expression of rat and human purinergic receptor P2X7 cDNA
      Buell, Gary Nutter; Surprenant, Annmarie; Kawashima, Eric
. IN
PA
      Glaxo Group Limited, UK
     U.S., 40 pp.
CODEN: USXXAM
SO
DТ
      Patent
LΑ
      English
 IC
      ICM C07H021-00
      ICS
          C07K014-475
      536023500
      3-2 (Biochemical Genetics)
 CC
      Section cross-reference(s): 13, 63
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                                            APPLICATION NO.
                                                                   DATE
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    US 6133434
                                20001017
                                            US 1997-842079
                                                                   19970428 <--
    US 6509163
                         В1
                                20030121
                                          US 2000-638857
                                                                   20000815 <--
PRAI US 1997-842079
                               19970428 <--
                         A1
CLASS
                CLASS PATENT FAMILY CLASSIFICATION CODES
PATENT NO.
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                       C07H021-00
US 6133434
                ICS
                       C07K014-475
                       536023500
                NCL
US 6509163
                ECLA
                       C07K014/705
   The present invention relates, in general, to a purinergic receptor and,
    in particular, to a P2X7 (also designated P2Z) receptor. Rat and human
    purinergic receptor P2X7 cDNA are cloned and expressed, and their tissue
    distribution and pharmacol. profiles are demonstrated. The invention also
    relates to a method of screening compds. for their ability to inhibit P2X7
    activity and thereby for their usefulness in treating a variety of
    diseases/disorders, including arthritic and respiratory disorders and
    neurodegenerative diseases.
ST
    rat human purinoceptor P2X cDNA sequence therapy; ATP receptor P2X7 cDNA
     sequence human rat
IT
    Purinoceptors
    RL: BOC (Biological occurrence); BPN (Biosynthetic preparation); BPR
     (Biological process); BSU (Biological study, unclassified); PRP
     (Properties); THU (Therapeutic use); BIOL (Biological study); OCCU
     (Occurrence); PREP (Preparation); PROC (Process); USES (Uses)
        (P2Z, of human and rat; cloning and expression of rat and human
       purinergic receptor P2X7 cDNA)
    Arthritis
TΤ
    Drug screening
    Gene therapy
    Genetic vectors
    Molecular cloning
        (cloning and expression of rat and human purinergic receptor P2X7 cDNA)
IT
    Antibodies
     RL: BPN (Biosynthetic preparation); THU (Therapeutic use); BIOL
     (Biological study); PREP (Preparation); USES (Uses)
        (cloning and expression of rat and human purinergic receptor P2X7 cDNA)
IT
    Nervous system
        (degeneration; cloning and expression of rat and human purinergic
       receptor P2X7 cDNA)
    Nucleic acid hybridization
TΤ
        (detection by; cloning and expression of rat and human purinergic
        receptor P2X7 cDNA)
IT
    Respiratory tract
        (disease; cloning and expression of rat and human purinergic receptor
        P2X7 cDNA)
IT
    Gene, animal
    RL: BPR (Biological process); BSU (Biological study, unclassified); PRP
     (Properties); THU (Therapeutic use); BIOL (Biological study); PROC
     (Process); USES (Uses)
        (for P2X7, of human and rat; cloning and expression of rat and human
       purinergic receptor P2X7 cDNA)
IT
    cDNA sequences
        (for human and rat purinoceptors P2X7; cloning and expression of rat
       and human purinergic receptor P2X7 cDNA)
TТ
    Diagnosis
        (mol.; cloning and expression of rat and human purinergic receptor P2X7
       cDNA)
TΤ
    Protein sequences
        (of human and rat purinoceptors P2X7; cloning and expression of rat and
       human purinergic receptor P2X7 cDNA)
IT
    159521-88-7
    RL: PRP (Properties)
        (Unclaimed; cloning and expression of rat and human purinergic receptor
    176521-57-6P, Receptor, purinergic P2x7 (rat brain)
    Purinoceptor P2X7 (human) 301457-00-1P, Purinoceptor P2X7 (human)
    RL: BOC (Biological occurrence); BPN (Biosynthetic preparation); BPR
     (Biological process); BSU (Biological study, unclassified); PRP
     (Properties); THU (Therapeutic use); BIOL (Biological study); OCCU
     (Occurrence); PREP (Preparation); PROC (Process); USES (Uses)
        (amino acid sequence; cloning and expression of rat and human
       purinergic receptor P2X7 cDNA)
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                  301457-01-2
TТ
     RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
     (Biological study)
        (nucleotide sequence; cloning and expression of rat and human
        purinergic receptor P2X7 cDNA)
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     US6133434 SEQID: 3 unclaimed DNA 301457-05-6, 4: PN: US6133434 SEQID: 4
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        (unclaimed nucleotide sequence; cloning and expression of rat and human
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                                  301335-90-0
                                                 301335-91-1
     301335-93-3
                   301335-94-4
                                  301335-95-5
                                                 301335-96-6
     RL: PRP (Properties)
        (unclaimed sequence; cloning and expression of rat and human purinergic
        receptor P2X7 cDNA)
RE.CNT 20
              THERE ARE 20 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE
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(19) Valera; Receptors and Channels 1995, V3(4), P283 HCAPLUS
(20) Wiley; Br J Pharmacol 1994, V112, P946 HCAPLUS
     301457-02-3, 1: PN: US6133434 SEQID: 1 unclaimed DNA
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        (unclaimed nucleotide sequence; cloning and expression of rat and human
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     301457-02-3 HCAPLUS
     1: PN: US6133434 SEQID: 1 unclaimed DNA (9CI) (CA INDEX NAME)
CN
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
L27 ANSWER 20 OF 25 HCAPLUS COPYRIGHT 2004 ACS on STN
     2000:316649 HCAPLUS
ΑN
     132:333387
     Entered STN: 16 May 2000
    Recombinant multivalent M protein vaccine against Streptococcus
TI
    Dale, James B.; Lederer, James W.
IN
    University of Tennessee Research Corporation, USA
PA
     U.S., 62 pp., Cont.-in-part of U.S. Ser. No. 945,954, abandoned.
so
    CODEN: USXXAM
ÐΤ
    Patent
    English
T.A
IC
     ICM A61K039-09
NCL 424244100
    15-2 (Immunochemistry)
CC
FAN.CNT 2
     PATENT NO.
                          KIND DATE
                                             APPLICATION NO.
                                                                      DATE
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     US 6063386
                                 20000516
                                              US 1997-937271
                                                                      19970915 <--
     ES 2170075
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                                            ES 1993-922201
                                                                      19930915 <--
                          Т3
PRAI US 1992-945954
                                 19920916 <--
                          B2
CLASS
                 CLASS PATENT FAMILY CLASSIFICATION CODES
PATENT NO.
US 6063386
                 TCM
                        A61K039-09
                 NCL
                        424244100
                       C07K014/315
                 ECLA
    The authors disclose the preparation of chimeric matrix proteins derived from
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multiple serotypes of group A streptococci. The chimeric proteins are

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immunogenic and provoke opsonic antibodies in rabbits.
ST
    Streptococcus multivalent M protein vaccine
    Proteins, specific or class
IT
    RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (M12; of chimeric multivalent matrix protein construct as vaccine
        against group A streptococci)
IT
    Proteins, specific or class
    RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (M18; of chimeric multivalent matrix protein construct as vaccine
        against group A streptococci)
IT
    Proteins, specific or class
    RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (M19; of chimeric multivalent matrix protein construct as vaccine
        against group A streptococci)
TT
    Proteins, specific or class
    RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (M1; of chimeric multivalent matrix protein construct as vaccine
        against group A streptococci)
IT
    Proteins, specific or class
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (M24; of chimeric multivalent matrix protein construct as vaccine
        against group A streptococci)
IT
    Proteins, specific or class
    RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (M2; of chimeric multivalent matrix protein construct as vaccine
        against group A streptococci)
    Proteins, specific or class
IT
    RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (M3; of chimeric multivalent matrix protein construct as vaccine
        against group A streptococci)
    Proteins, specific or class
TT
    RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (M4; of chimeric multivalent matrix protein construct as vaccine
        against group A streptococci)
    Proteins, specific or class
IT
    RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (M5; of chimeric multivalent matrix protein construct as vaccine
        against group A streptococci)
    Proteins, specific or class
IT
    RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (M6; of chimeric multivalent matrix protein construct as vaccine
        against group A streptococci)
    Peptides, biological studies
    RL: BOC (Biological occurrence); BSU (Biological study, unclassified);
    BIOL (Biological study); OCCU (Occurrence)
        (as linkers in chimeric multivalent matrix protein construct as vaccine
        against group A streptococci)
    Streptococcus group A
IT
        (chimeric multivalent M proteins as vaccine against)
IT
    Rheumatic fever
        (chimeric multivalent M proteins of group A streptococci as vaccine
        against)
    Fusion proteins (chimeric proteins)
TΤ
    RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES
     (Uses)
        (chimeric multivalent matrix protein construct as vaccine against group
        A streptococci)
TΤ
    Antibodies
    RL: BOC (Biological occurrence); BSU (Biological study, unclassified);
    BIOL (Biological study); OCCU (Occurrence)
        (crossreacting; with heart tissue in relation to chimeric multivalent {\tt M}
        protein vaccine against group A streptococci)
IT
    DNA sequences
    Protein sequences
        (for chimeric multivalent matrix proteins from group A streptococci)
    Synthetic gene
    Synthetic gene
    RL: PRP (Properties)
        (microbial; for chimeric multivalent matrix proteins from group A
        streptococci)
IT
        (pharyngitis, streptococcal; chimeric multivalent M proteins of group A
        streptococci as vaccine against)
IT
    Heart, disease
        (rheumatic; chimeric multivalent M proteins of group A streptococci as
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vaccine against)
TT
     Vaccines
        (synthetic; against group A streptococci using chimeric multivalent M
        proteins)
IT
     Gene, microbial
     Gene, microbial
     RL: PRP (Properties)
        (synthetic; for chimeric multivalent matrix proteins from group A
        streptococci)
TT
     Opsonins
     RL: BOC (Biological occurrence); BSU (Biological study, unclassified);
     BIOL (Biological study); OCCU (Occurrence)
        (to M proteins of group A streptococci induced by chimeric multivalent
        constructs)
TT
     153-94-6, D-Tryptophan
                               157-06-2, D-Arginine
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     344-25-2, D-Proline 640-68-6, D-Valine
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         (of peptide linkers in chimeric multivalent matrix protein construct as
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              THERE ARE 85 CITED REFERENCES AVAILABLE FOR THIS RECORD
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    P390
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TI

Search done by Noble Jarrell

Recombinant adenovirus particles (AdDYS.beta.gal) containing sequences

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encoding full length dystrophin used to treat patients with muscular
     dystrophy
     Kochanek, Stefan; Caskey, C. Thomas; Mitani, Kohnosuke; Clemens, Paula R.
IN
PΑ
     Baylor College of Medicine, USA; Howard Hughes Medical Institute
so
    U.S., 22 pp.
     CODEN: USXXAM
DΤ
    Patent
     English
LΑ
    ICM A01N043-04
IC
     ICS C12N015-00
NCL
   514044000
     1-12 (Pharmacology)
     Section cross-reference(s): 3, 10, 14
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PRAI US 1995-488014
                                19950607 <--
CLASS
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                NCL
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                ECLA C07K014/47A2A; C12N015/86F
US 5985846
    The invention provides a method for treatment of muscular dystrophy using
     mol. genetics. The method involves infecting muscle cells of muscular
     dystrophy patients with a recombinant adenovirus particle (AdDYS.beta.gal)
     which has encapsidated a gene transfer vector (pAdDYS.beta.gal). The gene
     transfer vector pAdDYS.beta.gal contains (5' to 3' orientation): (a) an
    adenovirus inverted terminal repeat (ITR); (b) an adenovirus packaging signal; (c) cDNA sequence encoding dystrophin operably linked to a mouse
     muscle creatine kinase gene (MCK) promoter; (d) a second adenovirus
     packaging signal and (e) a second adenovirus ITR. The infection of the
     recombinant adenovirus particle AdDYS.beta.gal results in recombinant
     expression of dystrophin which corrects the cytoskeletal defects found in
     the muscular dystrophy patient. The invention also presented: (a)
     construction of gene transfer vector pAdDYS.beta.gal; (b) production of
     recombinant adenovirus AdDYS.beta.gal using the gene transfer vector;
     expression of dystrophin in myoblasts transformed with AdDYS.beta.gal
     virus particles and (d) in vivo expression of dystrophin in mdx mice
     injected with AdDYS.beta.gal virus particles. The invention postulated
     that it may be possible to accommodate up to 37-kb of foreign DNA into the
     defective adenovirus, making it possible to deliver multiple or large
     genes containing tissue-specific or inducible promoters.
     gene therapy muscular dystrophy recombinant adenovirus AdDYSbetagal;
     dystrophin gene therapy muscular dystrophy recombinant adenovirus
     AdDYSbetagal; signal packaging ITR adenovirus AdDYSbetagal muscular
     dystrophy gene therapy; inverted terminal repeat adenovirus AdDYSbetagal
     muscular dystrophy gene therapy
IT
     Cytoskeleton
        (correction of cytoskeletal defects; recombinant adenovirus particles
        (AdDYS.beta.gal) containing sequences encoding full length dystrophin,
        adenovirus ITRs and packaging signals used to treat patients with
        muscular dystrophy)
IT
    Gene, animal
     RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
     (Uses)
        (for dystrophin; recombinant adenovirus particles (AdDYS.beta.gal)
        containing sequences encoding full length dystrophin used to treat patients
        with muscular dystrophy)
    Gene, animal
IT
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (for muscle creatine kinase, promoter of; recombinant adenovirus
        particles (AdDYS.beta.gal) containing sequences encoding full length
        dystrophin linked to a mouse muscle creatine kinase promoter used to
        treat patients with muscular dystrophy)
TT
    Repetitive DNA
     RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
        (inverted, terminal; recombinant adenovirus particles (AdDYS.beta.gal)
        containing sequences encoding full length dystrophin and adenovirus
        internal terminal repeats used to treat patients with muscular
        dystrophy)
IT
     Promoter (genetic element)
    RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
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(Uses)
         (of mouse muscle creatine kinase gene; recombinant adenovirus particles
         (AdDYS.beta.gal) containing sequences encoding full length dystrophin
        linked to a mouse muscle creatine kinase promoter used to treat
        patients with muscular dystrophy)
IT
     Virus vectors
         (pAdDYS.beta.gal; recombinant adenovirus particles (AdDYS.beta.gal)
        containing sequences encoding full length dystrophin used to treat patients
        with muscular dystrophy)
     Genetic element
TΤ
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); BUU (Biological use, unclassified); BIOL (Biological
     study); USES (Uses)
         (packaging signal; recombinant adenovirus particles (AdDYS.beta.gal)
        containing sequences encoding full length dystrophin, adenovirus ITRs and
        packaging signals used to treat patients with muscular dystrophy)
     Human adenovirus 5
TΤ
     Mouse
     Muscular dystrophy
     Myoblast
         (recombinant adenovirus particles (AdDYS.beta.gal) containing sequences
        encoding full length dystrophin used to treat patients with muscular
        dystrophy)
IТ
     Dystrophin
     RL: BPN (Biosynthetic preparation); THU (Therapeutic use); BIOL
     (Biological study); PREP (Preparation); USES (Uses)
         (recombinant adenovirus particles (AdDYS.beta.gal) containing sequences
        encoding full length dystrophin used to treat patients with muscular
        dystrophy)
     Molecular cloning
IT
         (recombinant adenovirus particles (AdDYS.beta.gal) containing sequences
        encoding full length dystrophin, adenovirus ITRs and packaging signals
        used to treat patients with muscular dystrophy)
IT
     Gene therapy
         (recombinant adenovirus particles (AdDYS.beta.gal) used in gene therapy
        for muscular dystrophy)
IT
     Human adenovirus
         (recombinant, AdDYS.beta.gal; recombinant adenovirus particles
         (AdDYS.beta.gal) containing sequences encoding full length dystrophin used
        to treat patients with muscular dystrophy)
TТ
     9001-15-4, Creatine kinase
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
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        muscle creatine kinase promoter used to treat patients with muscular
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         (unclaimed nucleotide sequence; recombinant adenovirus particles
         (AdDYS.beta.gal) containing sequences encoding full length dystrophin used
        to treat patients with muscular dystrophy)
              THERE ARE 13 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE
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ED
     Entered STN: 11 Nov 1999
    Human and mouse CD40 ligands and analogs and cDNAs encoding them and their
ΤI
    preparation and use
IN
     Armitage, Richard J.; Fanslow, William C.; Spriggs, Melanie K.;
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    Jeffrey T.
PA
    Immunex Corporation, USA
    U.S., 65 pp., Cont.-in-part of U.S. Ser. No. 249,189.
SO
     CODEN: USXXAM
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    English
LA
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    ICM C12N015-12
     ICS C12N015-63
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    536023500
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ECLA C07K014/705Q; C12N015/62
 US 6410711
     Membrane-bound and soluble forms of the cytokine CD40 ligand (CD40-L) of
     mouse and human are described and cDNAs encoding them are cloned and
     expressed. Expression constructs and host cells for manufacture of the protein
     are described. Deletion and substitution analogs of the ligand that retain specific binding to the extracellular binding region of a CD40
     receptor are described. Antisense nucleic acids and antibodies to the
     protein that can be used to modulate its action are described. Cell lines
     with high levels of CD40-L were screened for using a CD40 fusion protein
     with Ig/Fc as an affinity label and the mouse thymoma line EL-4 identified
     as high in the ligand. An expression cDNA library in the mammalian
     expression vector pDC406 was screened using the fusion protein upon
     expression in CV1-EBNA cells. The mouse clone was used as a probe to
     obtain a clone from a human peripheral blood lymphocyte cDNA library in
     .lambda.gt10. CD40-L stimulates B cell proliferation, the synthesis of
     polyclonal Igs and the shedding of Fc.epsilon.RII (CD23 antigen) from B
     cells.
ST
     CD40 ligand cDNA cloning expression; B cell proliferation CD40 ligand;
     cytotoxic T cell CD40 ligand
TT
     Immunoglobulins
     RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL
     (Biological study); PROC (Process)
        (A, CD40 ligand stimulation of production of; human and mouse CD40 ligands
        and analogs and cDNAs encoding them and their preparation and use)
ΙT
     Cell proliferation
        (B cell, CD40 ligand stimulation of; human and mouse CD40 ligands and
        analogs and cDNAs encoding them and their preparation and use)
TT
     Allergy inhibitors
        (CD40 derivs. as; human and mouse CD40 ligands and analogs and cDNAs
        encoding them and their preparation and use)
     Immunoglobulins
TΤ
     RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL
     (Biological study); PROC (Process)
        (CD40 ligand stimulation of production of; human and mouse CD40 ligands and
        analogs and cDNAs encoding them and their preparation and use)
     Glycoproteins, specific or class
TT
     RL: BPN (Biosynthetic preparation); PRP (Properties); THU (Therapeutic
     use); BIOL (Biological study); PREP (Preparation); USES (Uses)
        (CD40-L (antigen CD40 ligand); human and mouse CD40 ligands and analogs
        and cDNAs encoding them and their preparation and use)
IT
     Immunoglobulins
     RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL
     (Biological study); PROC (Process)
        (E, CD40 ligand stimulation of production of; human and mouse CD40 ligands and analogs and cDNAs encoding them and their preparation and use)
     Immunoglobulins
     RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL
     (Biological study); PROC (Process)
        (G1, CD40 ligand stimulation of production of; human and mouse CD40 ligands
        and analogs and cDNAs encoding them and their preparation and use)
     Immunoglobulins
     RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL
     (Biological study); PROC (Process)
        (G2b, CD40 ligand stimulation of production of; human and mouse CD40
        ligands and analogs and cDNAs encoding them and their preparation and use)
     Immunoglobulins
     RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL
     (Biological study); PROC (Process)
        (G3, CD40 ligand stimulation of production of; human and mouse CD40 ligands
        and analogs and cDNAs encoding them and their preparation and use)
     Immunoglobulin receptors
     RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL (Biological study); PROC (Process)
        (IgE type II, CD40 ligand stimulation of shedding of; human and mouse
        CD40 ligands and analogs and cDNAs encoding them and their preparation and
        use)
IT
     Immunoglobulins
     RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL
     (Biological study); PROC (Process)
        (M, CD40 ligand stimulation of production of; human and mouse CD40 ligands and analogs and cDNAs encoding them and their preparation and use)
IT
     Gene, animal
     RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
     (Biological study)
        (cDNA, for CD40 ligand of mouse and human; human and mouse CD40 ligands
        and analogs and cDNAs encoding them and their preparation and use)
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TT
    Immunity
        (cell-mediated, CD40-L and stimulation of; human and mouse CD40 ligands
        and analogs and cDNAs encoding them and their preparation and use)
TT
    T cell (lymphocyte)
        (cytotoxic, CD40-L stimulation of proliferation of; human and mouse
        CD40 ligands and analogs and cDNAs encoding them and their preparation and
        use)
IT
    cDNA sequences
        (for CD40 ligand of human and mouse; human and mouse CD40 ligands and
        analogs and cDNAs encoding them and their preparation and use)
IT
    Immunoglobulins
    RL: BPN (Biosynthetic preparation); BIOL (Biological study); PREP
     (Preparation)
        (fusion products, with CD40 ligand, preparation of; human and mouse CD40
        ligands and analogs and cDNAs encoding them and their preparation and use)
        (human and mouse CD40 ligands and analogs and cDNAs encoding them and
        their preparation and use)
IT
    Antibodies
    RL: ARG (Analytical reagent use); BSU (Biological study, unclassified);
    THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study);
    USES (Uses)
        (monoclonal, to CD40 ligand; human and mouse CD40 ligands and analogs
        and cDNAs encoding them and their preparation and use)
IT
    Protein sequences
        (of CD40 ligand of human and mouse; human and mouse CD40 ligands and
        analogs and cDNAs encoding them and their preparation and use)
IT
    Antibodies
    RL: ARG (Analytical reagent use); BSU (Biological study, unclassified);
    THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study);
    USES (Uses)
        (to CD40 ligand; human and mouse CD40 ligands and analogs and cDNAs
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     (Biological study); PROC (Process)
        (amino acid sequence; human and mouse CD40 ligands and analogs and
        cDNAs encoding them and their preparation and use)
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    157214-04-5DP, fusion products with CD40 ligand
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        (amino acid sequence; human and mouse CD40 ligands and analogs and
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(2) Anon; EP 0330191 1989 HCAPLUS
(3) Anon; EP 0555880 1993 HCAPLUS
(4) Anon; WO 9309812 1993 HCAPLUS
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(29) Paulie; J Immunol 1989, V142, P590 HCAPLUS
(30) Potocnik; Scan J Immunol 1990, V31, P213 MEDLINE
(31) Rousset; J Exp Med 1991, V173, P705 HCAPLUS (32) Spriggs; J Exp Med 1992, V176, P1543 HCAPLUS
(33) Srinivasan; US 5716805 1998 HCAPLUS
(34) Stamenkovic; EMBO J 1989, V8, P1403 HCAPLUS
(35) Valle; Eur J Immunol 1989, V19, P1463 HCAPLUS
(36) Wang; J Virol 1991, V65, P4101 HCAPLUS
(37) Zettlmeissl; DNA Cell Biol 1990, V9, P347 HCAPLUS
(38) Zhang; J Immunol 1991, V146, P1836 HCAPLUS
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ED
     Recombinant soluble CD40 ligand polypeptide and pharmaceutical composition
TI
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     Armitage, Richard J.; Fanslow, William C.; Spriggs, Melanie K.;
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     Srinivasan, Subhashini; Gibson, Marylou G.; Morris, Arvia E.; McGrew,
     Jeffrey T.
     Immunex Corporation, USA
PA
     U.S., 64 pp., Cont.-in-part of U.S. Ser. No. 249,189. CODEN: USXXAM
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     15-2 (Immunochemistry)
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    Disclosed are polypeptides (e.g. membrane bound CD40-L, monomeric and oligomeric CD40-L, soluble CD40-L, and fusion protein of CD40-L) and
     antisense and sense DNA and RNA sequences, vectors and transformed host
     cells useful in providing CD40-L polypeptides. More particularly, this
     invention provides isolated human and murine CD40-L polypeptides that bind
     to the extracellular binding region of a CD40 receptor. Also, provided
     are soluble CD40 comprising extracellular region of human CD40 and CD40/Fc
     fusion proteins, and antibodies. CD40 agonists, i.e. membrane-bound
     CD40-L and oligomeric CD40-L, are useful as vaccine adjuvant and for
     stimulating B cell proliferation and monoclonal antibody production from
     hybridoma cells. CD40 antagonists, i.e. CD40 receptor, CD40/Fc fusion
     protein, soluble CD40/Fc, monomeric CD40-L, are useful for treating
     autoimmune diseases characterized by presence of high levels of
     antigen-antibody complexes, such as allergy, insulin dependent diabetes
     mellitus, graft vs. host disease and others.
     CD40 ligand oligomer fusion protein; soluble CD40 ligand antisense DNA
ST
     RNA; monoclonal antibody CD40 ligand adjuvant; autoimmune disease B cell
     proliferation antiallergic
TТ
     Cell proliferation
        (B cell; recombinant soluble CD40 ligand polypeptide as CD40 agonist and
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antagonist as adjuvant, stimulant of B cell proliferation and antibody
        production, and for treating autoimmune diseases)
     Glycoproteins, specific or class
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (CD40-L (antigen CD40 ligand); recombinant soluble CD40 ligand polypeptide
        as CD40 agonist and antagonist as adjuvant, stimulant of B cell proliferation and antibody production, and for treating autoimmune
        diseases)
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     RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL
     (Biological study); PROC (Process)
        (CD40; recombinant soluble CD40 ligand polypeptide as CD40 agonist and
        antagonist as adjuvant, stimulant of B cell proliferation and antibody
        production, and for treating autoimmune diseases)
IT
     Immunoglobulins
     RL: BOC (Biological occurrence); BSU (Biological study, unclassified);
     BIOL (Biological study); OCCU (Occurrence)
        (E, secretion induction; recombinant soluble CD40 ligand polypeptide as
        CD40 agonist and antagonist as adjuvant, stimulant of B cell
        proliferation and antibody production, and for treating autoimmune
        diseases)
     Animal cell line
IT
        (EL4; recombinant soluble CD40 ligand polypeptide as CD40 agonist and
        antagonist as adjuvant, stimulant of B cell proliferation and antibody
        production, and for treating autoimmune diseases)
IT
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES
     (Uses)
        (G1, fragment fusion protein; recombinant soluble CD40 ligand polypeptide
        as CD40 agonist and antagonist as adjuvant, stimulant of B cell
        proliferation and antibody production, and for treating autoimmune
        diseases)
IT
     Immunoglobulins
     RL: BOC (Biological occurrence); BSU (Biological study, unclassified); THU
     (Therapeutic use); BIOL (Biological study); OCCU (Occurrence); USES (Uses)
        (G; recombinant soluble CD40 ligand polypeptide as CD40 agonist and antagonist as adjuvant, stimulant of B cell proliferation and antibody
        production, and for treating autoimmune diseases)
     Immunoglobulin receptors
TT
     RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL
     (Biological study); PROC (Process)
        (IgE type II; recombinant soluble CD40 ligand polypeptide as CD40 agonist
        and antagonist as adjuvant, stimulant of B cell proliferation and
        antibody production, and for treating autoimmune diseases)
IT
     Immunoglobulins
     RL: BOC (Biological occurrence); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); OCCU (Occurrence); USES (Uses)
        (M; recombinant soluble CD40 ligand polypeptide as CD40 agonist and
        antagonist as adjuvant, stimulant of B cell proliferation and antibody
        production, and for treating autoimmune diseases)
     Antigens
IT
     RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL
     (Biological study); PROC (Process)
        (T-cell-dependent antigens; recombinant soluble CD40 ligand polypeptide as
        CD40 agonist and antagonist as adjuvant, stimulant of B cell
        proliferation and antibody production, and for treating autoimmune
        diseases)
IT
     Immunostimulants
        (adjuvants; recombinant soluble CD40 ligand polypeptide as CD40 agonist
        and antagonist as adjuvant, stimulant of B cell proliferation and
        antibody production, and for treating autoimmune diseases)
IT
     Drug delivery systems
        (carriers; recombinant soluble CD40 ligand polypeptide as CD40 agonist and
        antagonist as adjuvant, stimulant of B cell proliferation and antibody
        production, and for treating autoimmune diseases)
IT
    T cell (lymphocyte)
        (cytotoxic, induction; recombinant soluble CD40 ligand polypeptide as CD40
        agonist and antagonist as adjuvant, stimulant of B cell proliferation
        and antibody production, and for treating autoimmune diseases)
IT
    Immunoassav
        (enzyme-linked immunosorbent assay; recombinant soluble CD40 ligand
        polypeptide as CD40 agonist and antagonist as adjuvant, stimulant of B
        cell proliferation and antibody production, and for treating autoimmune
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diseases)
TT
     Transplant and Transplantation
        (graft-vs.-host reaction; recombinant soluble CD40 ligand polypeptide as
        CD40 agonist and antagonist as adjuvant, stimulant of B cell
        proliferation and antibody production, and for treating autoimmune
        diseases)
IT'
     T cell (lymphocyte)
        (helper cell; recombinant soluble CD40 ligand polypeptide as CD40 agonist
        and antagonist as adjuvant, stimulant of B cell proliferation and
        antibody production, and for treating autoimmune diseases)
IT
        (immunoblotting, dot; recombinant soluble CD40 ligand polypeptide as CD40
        agonist and antagonist as adjuvant, stimulant of B cell proliferation
        and antibody production, and for treating autoimmune diseases)
IT
     Diabetes mellitus
        (insulin-dependent; recombinant soluble CD40 ligand polypeptide as CD40
        agonist and antagonist as adjuvant, stimulant of B cell proliferation
        and antibody production, and for treating autoimmune diseases)
TT
     Antibodies
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES
     (Uses)
        (monoclonal; recombinant soluble CD40 ligand polypeptide as CD40 agonist
        and antagonist as adjuvant, stimulant of B cell proliferation and
        antibody production, and for treating autoimmune diseases)
IT
     Mononuclear cell (leukocyte)
        (peripheral blood; recombinant soluble CD40 ligand polypeptide as CD40
        agonist and antagonist as adjuvant, stimulant of B cell proliferation
        and antibody production, and for treating autoimmune diseases)
IT
    Blood
        (peripheral; recombinant soluble CD40 ligand polypeptide as CD40 agonist
        and antagonist as adjuvant, stimulant of B cell proliferation and
        antibody production, and for treating autoimmune diseases)
IT
     Immunoglobulins
     RL: BOC (Biological occurrence); BPN (Biosynthetic preparation); BSU
     (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological
     study); OCCU (Occurrence); PREP (Preparation); USES (Uses)
        (polyclonal; recombinant soluble CD40 ligand polypeptide as CD40 agonist
        and antagonist as adjuvant, stimulant of B cell proliferation and
        antibody production, and for treating autoimmune diseases)
    Allergy
     Allergy inhibitors
Autoimmune disease
     Hybridoma
     Molecular cloning
     Protein sequences
     Vaccines
     cDNA library
     cDNA sequences
        (recombinant soluble CD40 ligand polypeptide as CD40 agonist and
        antagonist as adjuvant, stimulant of B cell proliferation and antibody
        production, and for treating autoimmune diseases)
IT
     Interleukin 4
     RL: BAC (Biological activity or effector, except adverse); BPR (Biological
     process); BSU (Biological study, unclassified); BIOL (Biological study);
     PROC (Process)
        (recombinant soluble CD40 ligand polypeptide as CD40 agonist and
        antagonist as adjuvant, stimulant of B cell proliferation and antibody
        production, and for treating autoimmune diseases)
    Antibodies
     RL: BOC (Biological occurrence); BPN (Biosynthetic preparation); BSU
     (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological
     study); OCCU (Occurrence); PREP (Preparation); USES (Uses)
        (recombinant soluble CD40 ligand polypeptide as CD40 agonist and
        antagonist as adjuvant, stimulant of B cell proliferation and antibody production, and for treating autoimmune diseases)
IT
    CD40 (antigen)
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (recombinant soluble CD40 ligand polypeptide as CD40 agonist and
        antagonist as adjuvant, stimulant of B cell proliferation and antibody
        production, and for treating autoimmune diseases)
    Fusion proteins (chimeric proteins)
IT
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP
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(Preparation); USES (Uses)
        (recombinant soluble CD40 ligand polypeptide as CD40 agonist and
        antagonist as adjuvant, stimulant of B cell proliferation and antibody
        production, and for treating autoimmune diseases)
     Antisense DNA
IT
     Antisense RNA
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES
     (Uses)
        (recombinant soluble CD40 ligand polypeptide as CD40 agonist and
        antagonist as adjuvant, stimulant of B cell proliferation and antibody
        production, and for treating autoimmune diseases)
ΙT
     DNA
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     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES
     (Uses)
        (sense; recombinant soluble CD40 ligand polypeptide as CD40 agonist and
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        production, and for treating autoimmune diseases)
     148814-08-8, CD40 ligand (human) 149119-89-1 155979-32-1 186361-42-2, CD40 ligand [194-tryptophan] (human) 186361-47-7
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     245524-10-1 245524-52-1, 119-260-CD40 liqand (mouse) 245524-56-5,
     51-261-CD40 ligand (human clone CD40-L)
     RL: PRP (Properties)
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RE
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(2) Anon; EP 0330191 1989 HCAPLUS
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(45) Zhang; J Immunol 1991, V146, P1836 HCAPLUS
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     comprising the same and hybridomas producing the same
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     Immunex Corporation, USA
     U.S., 59 pp., Cont.-in-part of U.S. Ser. No. 969,703, abandoned. .
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     ICM C07K016-28
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US 6391637
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US 2003091564
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US 6410711
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                       C07K014/705Q; C12N015/62
US 2003144182
                 ECLA
AB Disclosed is a polypeptide (CD40-L) and DNA sequences, vectors and
     transformed host cells useful in providing CD40-L polypeptides. More
     particularly, this invention provides isolated human and murine CD40-L
     polypeptides that bind to the extracellular binding region of a CD40
     receptor. The invention further provides CD40-L fragment for production of
     monoclonal antibodies specific for CD40-L. Also, soluble CD40 protein and
     CD40 fusion proteins are prepared for inducing B cell proliferation and
     antibody (e..g IgE) secretion, and for anti-allergic treatment. Also,
     fusion proteins comprising soluble human or murine CD40-L and Fc or trimeric
     CD40-L are constructed. In summary, CD40 agonists (i.e. membrane-bound
     CD40-L and oligomeric CD40-L) are provided for use as vaccine adjuvant and
     antibody production stimulant, and CD40 antagonists (i.e. CD40 receptor,
     CD40/Fc, and soluble monomeric CD40-L) are provided for treating autoimmune
ST
     CD40 ligand fusion protein monoclonal antibody
     Cell proliferation
IT
        (B cell; monoclonal antibodies to CD40 ligand, pharmaceutical composition
        comprising the same and hybridomas producing the same)
IT
        (CD40 ligand fragments of mouse and human in production of monoclonal antibodies to CD40 ligand and preparation of CD40 agonist and antagonists
        and their therapeutical applications)
     Glycoproteins, specific or class
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified);
     PRP (Properties); PUR (Purification or recovery); BIOL (Biological study);
     PREP (Preparation)
        (CD40-L (antigen CD40 ligand); monoclonal antibodies to CD40 ligand,
        pharmaceutical composition comprising the same and hybridomas producing the
        same)
TT
     Receptors
     RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL
     (Biological study); PROC (Process)
        (CD40; monoclonal antibodies to CD40 ligand, pharmaceutical composition
        comprising the same and hybridomas producing the same)
IT
    Immunoglobulins
     RL: BOC (Biological occurrence); BPR (Biological process); BSU (Biological
     study, unclassified); BIOL (Biological study); OCCU (Occurrence); PROC
     (Process)
        (E, secretion; monoclonal antibodies to CD40 ligand, pharmaceutical
        composition comprising the same and hybridomas producing the same)
IT
        (EL4; monoclonal antibodies to CD40 ligand, pharmaceutical composition
        comprising the same and hybridomas producing the same)
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IT

Immunoglobulins

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RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (G1; monoclonal antibodies to CD40 ligand, pharmaceutical composition
        comprising the same and hybridomas producing the same)
     Secretion (process)
        (Ig; monoclonal antibodies to CD40 ligand, pharmaceutical composition
        comprising the same and hybridomas producing the same)
TT
     Immunoglobulin receptors
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (IGE type II; monoclonal antibodies to CD40 ligand, pharmaceutical
        composition comprising the same and hybridomas producing the same)
IT
     Immunostimulants
        (adjuvants; monoclonal antibodies to CD40 ligand, pharmaceutical composition
        comprising the same and hybridomas producing the same)
IT
    Drug delivery systems
        (carriers; monoclonal antibodies to CD40 ligand, pharmaceutical composition
        comprising the same and hybridomas producing the same)
TТ
     Allergy inhibitors
     Autoimmune disease
     Drug delivery systems
     Hybridoma
     Immunostimulants
     Molecular cloning
     Protein sequences
     Vaccines
     cDNA sequences
        (monoclonal antibodies to CD40 ligand, pharmaceutical composition comprising
        the same and hybridomas producing the same)
TТ
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); BIOL (Biological study)
        (monoclonal antibodies to CD40 ligand, pharmaceutical composition comprising
        the same and hybridomas producing the same)
     Fusion proteins (chimeric proteins)
     RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (monoclonal antibodies to CD40 ligand, pharmaceutical composition comprising
        the same and hybridomas producing the same)
ΤТ
     Antibodies
     RL: BPN (Biosynthetic preparation); THU (Therapeutic use); BIOL
     (Biological study); PREP (Preparation); USES (Uses)
        (monoclonal antibodies to CD40 ligand, pharmaceutical composition comprising
        the same and hybridomas producing the same)
TT
     Antisense DNA
     Antisense RNA
     DNA
     RNA
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (monoclonal antibodies to CD40 ligand, pharmaceutical composition comprising
        the same and hybridomas producing the same)
     CD40 (antigen)
     RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (monoclonal antibodies to CD40 ligand, pharmaceutical composition comprising
        the same and hybridomas producing the same)
     Antibodies
     RL: BPN (Biosynthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
        (monoclonal; monoclonal antibodies to CD40 ligand, pharmaceutical
        composition comprising the same and hybridomas producing the same)
     Immunoglobulins
     RL: BSU (Biological study, unclassified); MFM (Metabolic formation); THU
     (Therapeutic use); BIOL (Biological study); FORM (Formation,
     nonpreparative); USES (Uses)
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     148814-08-8, CD40 ligand (human) 149119-89-1 155979-32-1
IT
                                                                      186361-47-7
                   245524-07-6, 47-260-CD40 ligand (mouse)
     245524-02-1
                                                              245524-10-1
     RL: PRP (Properties)
        (amino acid sequence; CD40 ligand fragments of mouse and human in
        production of monoclonal antibodies to CD40 ligand and preparation of CD40
        agonist and antagonists and their therapeutical applications)
     149119-87-9, DNA (human CD40 ligand cDNA plus flanks)
     212581-01-6, DNA (mouse CD40 ligand cDNA)
                                                 245524-01-0 245524-04-3
     RL: PRP (Properties)
        (nucleotide sequence; CD40 ligand fragments of mouse and human in
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production of monoclonal antibodies to CD40 ligand and preparation of CD40
        agonist and antagonists and their therapeutical applications)
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     245525-96-6, PN: US5962406 SEQID: 14 unclaimed DNA 245525-99-9, PN:
     US5962406 SEOID: 18 unclaimed DNA 245526-07-2, PN: US5961974 SEQID: 19
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     RL: PRP (Properties)
        (unclaimed nucleotide sequence; monoclonal antibodies to CD40 ligand,
        pharmaceutical composition comprising the same and hybridomas producing the
        same)
TT
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     RL: PRP (Properties)
        (unclaimed protein sequence; monoclonal antibodies to CD40 ligand,
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     RL: PRP (Properties)
        (unclaimed protein sequence; monoclonal antibodies to CD40 ligand,
        pharmaceutical composition comprising the same and hybridomas producing the
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(42) Zhang; J Immunol 1991, V146, P1836 HCAPLUS
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        (unclaimed nucleotide sequence; monoclonal antibodies to CD40 ligand,
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    129:201670
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    Entered STN: 15 Sep 1998
ED
    Somatic mutations in the multiple tumor suppressor gene MTS1 and their use
TI
     in the diagnosis and prognosis of cancers
IN
     Kamb, Alexander
PΑ
    Myriad Genetics, Inc., USA
    U.S., 73 pp., Cont.-in-part of U.S. Ser. No. 251,938, abandoned.
SO
     CODEN: USXXAM
DT
     Patent
     English
LΑ
    ICM C07H021-04
IC
     ICS C12N015-63
NCL
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    14-1 (Mammalian Pathological Biochemistry)
CC
     Section cross-reference(s): 3, 9
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                 CLASS PATENT FAMILY CLASSIFICATION CODES
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                 ICS
                 NCL
                        536024310
AB
    The present invention relates to somatic mutations in the Multiple Tumor
     Suppressor (MTS) gene in human cancers and their use in the diagnosis and
    prognosis of human cancer. The invention further relates to germ line
     mutations in the MTS gene and their use in the diagnosis of predisposition
     to melanoma, leukemia, astrocytoma, glioblastoma, lymphoma, glioma,
     Hodgkin's lymphoma, CLL, and cancers of the pancreas, breast, thyroid,
    ovary, uterus, testis, kidney, stomach and rectum. The invention also relates to the therapy of human cancers which have a mutation in the MTS
     gene, including gene therapy, protein replacement therapy and protein
     mimetics. Finally, the invention relates to the screening of drugs for
     cancer therapy. Map-based cloning of the gene is described.
ST
    multiple tumor suppressor gene MTS1 human; somatic mutation MTS1 gene
     cancer
IT
    Gene, animal
     RL: BPR (Biological process); BSU (Biological study, unclassified); PRP
     (Properties); THU (Therapeutic use); BIOL (Biological study); PROC
     (Process); USES (Uses)
        (MTS2; somatic mutations in multiple tumor suppressor gene MTS1 and
        their use in diagnosis and prognosis of cancers)
IT
    Astrocyte
        (astrocytoma, susceptibility, diagnosis, and prognosis; somatic
        mutations in multiple tumor suppressor gene MTS1 and their use in
        diagnosis and prognosis of cancers)
IT
        (chronic lymphocytic, susceptibility, diagnosis, and prognosis; somatic
        mutations in multiple tumor suppressor gene MTS1 and their use in
        diagnosis and prognosis of cancers)
IT
     Intestine, neoplasm
        (colorectal, susceptibility, diagnosis, and prognosis; somatic
       mutations in multiple tumor suppressor gene MTS1 and their use in diagnosis and prognosis of cancers)
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245525-84-2 HCAPLUS

RN

CN

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Primers (nucleic acid)
    RL: ARG (Analytical reagent use); THU (Therapeutic use); ANST (Analytical
    study); BIOL (Biological study); USES (Uses)
        (for detection of somatic mutation in MTS1 gene; somatic mutations in
       multiple tumor suppressor gene MTS1 and their use in diagnosis and
       prognosis of cancers)
TT
    Neuroglia
        (glioblastoma, susceptibility, diagnosis, and prognosis; somatic
       mutations in multiple tumor suppressor gene MTS1 and their use in
       diagnosis and prognosis of cancers)
IT
    Neuroglia
        (glioma, susceptibility, diagnosis, and prognosis; somatic mutations in
       multiple tumor suppressor gene MTS1 and their use in diagnosis and
       prognosis of cancers)
    Chromosome
TT
        (human 9, 9p-ter, MTS1 and MTS2 genes mapping to; somatic mutations in
       multiple tumor suppressor gene MTS1 and their use in diagnosis and
       prognosis of cancers)
TT
    Diagnosis
        (mol.; somatic mutations in multiple tumor suppressor gene MTS1 and
       their use in diagnosis and prognosis of cancers)
TT
    Gene, animal
    RL: BPR (Biological process); BSU (Biological study, unclassified); PRP
     (Properties); THU (Therapeutic use); BIOL (Biological study); PROC
     (Process); USES (Uses)
        (mtsl; somatic mutations in multiple tumor suppressor gene MTSl and
       their use in diagnosis and prognosis of cancers)
TT
    Mammary gland
        (neoplasm, susceptibility, diagnosis, and prognosis; somatic mutations
        in multiple tumor suppressor gene MTS1 and their use in diagnosis and
       prognosis of cancers)
TT
    DNA sequences
        (of MTS1 and MTS2 genes of human; somatic mutations in multiple tumor
        suppressor gene MTS1 and their use in diagnosis and prognosis of
       cancers)
    Protein sequences
IT
        (of MTS1 gene product of human; somatic mutations in multiple tumor
        suppressor gene MTS1 and their use in diagnosis and prognosis of
       cancers)
    Genetic mapping
IT
        (of MTS1 gene; somatic mutations in multiple tumor suppressor gene MTS1
        and their use in diagnosis and prognosis of cancers)
IT
        (of MTS1-dependent tumors; somatic mutations in multiple tumor
        suppressor gene MTS1 and their use in diagnosis and prognosis of
        cancers)
    Susceptibility (genetic)
IT
        (somatic mutations in multiple tumor suppressor gene MTS1 and their use
        in diagnosis and prognosis of cancers)
IT
    Hodgkin's disease
    Kidney, neoplasm
    Leukemia
    Lymphoma
    Melanoma
    Ovary, neoplasm
     Pancreas, neoplasm
     Stomach, neoplasm
    Testis, neoplasm
    Thyroid gland, neoplasm
    Uterus, neoplasm
        (susceptibility, diagnosis, and prognosis; somatic mutations in
       multiple tumor suppressor gene MTS1 and their use in diagnosis and
       prognosis of cancers)
TT
    153520-02-6
    RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
     (Biological study)
        (amino acid sequence; somatic mutations in multiple tumor suppressor
       gene MTS1 and their use in diagnosis and prognosis of cancers)
    211994-88-6
IT
    RL: BSU (Biological study, unclassified); PRP (Properties); BIOL
     (Biological study)
        (nucleotide sequence; somatic mutations in multiple tumor suppressor
       gene MTS1 and their use in diagnosis and prognosis of cancers)
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    211994-69-3
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RL: BSU (Biological study, unclassified); PRP (Properties); THU
     (Therapeutic use); BIOL (Biological study); USES (Uses)
         (nucleotide sequence; somatic mutations in multiple tumor suppressor
        gene MTS1 and their use in diagnosis and prognosis of cancers)
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     (Therapeutic use); BIOL (Biological study); USES (Uses)
         (nucleotide sequence; somatic mutations in multiple tumor suppressor
        gene MTS1 and their use in diagnosis and prognosis of cancers)
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